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(71) Applicant (for all designated States except US): THE GOVERNORS OF THE UNIVERSITY OF ALBERTA [CA/CA]; 4000 RESEARCH TRANSITION FACILITY(RTF), 8308-114 Street NW, Edmonton, Alberta T6G 2E1 (CA).

## (72) Inventors; and

(75) Inventors/Applicants (for US only): KNAUS, Edward, E. [CA/CA]; 3531-104A Street, Edmonton, Alberta T6J 2K2 (CA). VELAZQUEZ, Carlos [MX/US]; 1470 West Key Parkway, Frederick, MD 21702 (US).

(74) Agent: CHAN, Albert, Wai-kit; LAW OFFICES OF ALBERT WAI-KIT CHAN LLC, WORLD PLAZA, SUITE 604, 141-07 20th Avenue, Whitestone, NY 11357 (US).

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(54) Title: NOVEL NSAIDS POSSESSING A NITRIC OXIDE DONOR DIAZEN-1-IUM-1,2-DIOLATE MOIETY

(57) Abstract: This invention provides a prodrug that help arthritis patients without increasing cardiovascular and gastrointestinal risk. A novel group of hybrid nitric oxide-releasing non-steroidal anti-inflammatory drugs (NO-NSAIDs), moiety attached via a one -carbon methylene spacer to the carboxylic acid group of the traditional NSAIDs aspirin, ibuprofen and indomethacin were synthesized. The ester prodrugs showed equipotent anti-inflammatory activities in vivo to that of the parent aspirin, ibuprofen and indomethacin. The simultaneous release of parent drug and nitric oxide from the NO- prodrugs constitutes a potentially beneficial property for the prophylactic prevention of thrombus formation and adverse cardiovascular events such as stroke and myocardial infarction. Data acquired in an in vivo ulcer index (UI) assay showed that this group of ester prodrugs in which no lesions were observed when compared to the parent drugs at equivalent doses. Accordingly, these hybrid NO-NSAID prodrugs possessing a diazen-1-ium-1, 2-diolate moiety, represents a new approach for the rational design of anti-inflammatory drugs with reduced gastric ulcerogenicity.

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NOVEL NSAIDS POSSESSING A NITRIC OXIDE DONOR  
DIAZEN-1-IUM-1,2-DIOLATE MOIETY

5 This application claims the benefits of U.S. provisional application 60/728,364, filed October 19, 2005, and U.S. provisional application 60/681,842, filed May 16, 2005. The contents of these preceding applications are hereby incorporated in their entireties by reference into this application.

10

Throughout this application, various publications are referenced. The disclosures of these publications in their entireties are hereby incorporated by reference into this application to more fully describe the state of the art to which this invention pertains.

15

BACKGROUND OF THE INVENTION

20 This invention provides a prodrug that help arthritis patients without increasing cardiovascular and gastrointestinal risk.

The major mechanism of action by which non-steroidal anti-inflammatory drugs (NSAIDs) exhibit anti-inflammatory activity involves the inhibition of cyclooxygenase (COX)-derived prostaglandin (PG) synthesis.<sup>1-4</sup> PGs, in addition to being undesirable effectors of inflammatory reactions, also exert important physiological functions such as gastrointestinal cytoprotection and vascular homeostasis.<sup>5-7</sup> In this regard, drugs that are more selective inhibitors of the COX-2 isozyme, relative to the COX-1 isozyme, allow the beneficial synthesis of cytoprotective PGs in the stomach in conjunction with a simultaneous inhibition of proinflammatory PG synthesis in joints. Chronic use of NSAIDs is associated with alterations in gastrointestinal integrity and function<sup>8,9</sup> which results in

30

the development of gastric ulcers.<sup>10</sup> Thus, the gastric irritant effect of aspirin (1) can be a deterrent to its long-term use for the prophylactic prevention of adverse cardiovascular events such as stroke and myocardial infarction.<sup>11,12</sup> Aspirin is  
5 a unique nonselective COX inhibitor due to its ability to acetylate the Ser530 hydroxyl group in the primary COX binding site of COX-1 and COX-2. In this regard, aspirin is a 10- to 100-fold more potent inhibitor of COX-1 relative to COX-2.<sup>13</sup> Acetylation of the weakly nucleophilic OH of Ser530 by aspirin  
10 is thought to result from initial binding of its COOH to Arg120 near the mouth of the COX binding site, which positions the ortho-acetoxy moiety in close proximity to the Ser530 OH, which it acetylates. Orally administered aspirin irreversibly acetylates Ser530 of COX-1 in platelets,<sup>14</sup> which results in a  
15 complete inhibition of platelet-derived thromboxane A2 (TxA2) biosynthesis. TxA2 is a potent platelet aggregator which also induces vasoconstriction and smooth muscle proliferation.<sup>15,16</sup> However, there remains a significant risk of gastrointestinal bleeding<sup>17-19</sup> due to inhibition of COX-1-mediated gastric PG  
20 synthesis even with low prophylactic doses of aspirin.<sup>20-23</sup>

COX-2 inhibitors are new, and in many ways, an improved class of drugs that are designed to be equally effective as traditional NSAIDS but safer. Traditional NSAIDS such as  
25 aspirin, Motrin, Aleve and other prescription drugs act by blocking the production of a family of chemicals that cause inflammation known as prostaglandins. Two enzymes appear to be crucial for the production of these prostaglandins, namely COX-1 and COX-2. Traditional NSAIDS inhibit both COX-1 and  
30 COX-2. Unfortunately, this nonselective inhibition of both COX-1 and COX-2 also inhibits prostaglandins involved in helping blood to clot, and protecting our stomach from ulcers. It is now strongly believed that this non-selective inhibition of both COX-1 and COX-2 by aspirin and related compounds is

why NSAIDS carry a risk of bleeding and stomach ulcerations. A new class of drugs, namely the COX-2 inhibitors, only inhibits the enzyme involved in inflammation and leaves our physiologic housekeeping functions alone.

5

However, the safety of COX-2 inhibitors has been questioned. The most famous event is that a blockbuster drug from Merck Vioxx was pulled off from pharmacy shelves in 2004 after Merck's trials showed an increased risk of heart and stroke damage. The two other COX-2 inhibitors on the market Celebrex and Bextra, are under intense study for their safety. On April 7, 2005, the Food and Drug Administration requested that Pfizer suspend sales of Bextra in the United States. The Food and Drug Administration is requiring all prescription anti-inflammatory arthritis medicines to provide additional information about cardiovascular and gastrointestinal risk.

15

Nitric oxide (NO) is now widely recognized as a critical mediator of gastrointestinal mucosal defense, exerting many of the same actions as prostaglandins in the gastrointestinal tract.<sup>10</sup> NO has been shown to reduce the severity of gastric injury in experimental models.<sup>24,25</sup> It has been proposed that the linking of an NO-releasing moiety to an NSAID may reduce the toxicity of the latter.<sup>26</sup> In animal studies, NO-releasing derivatives of a wide range of NSAIDs (Figure 1) including the NO-aspirin (2), NO-naproxen (3), NO-flurbiprofen (4) and NO-diclofenac (5), have been shown to spare the gastrointestinal tract, even though they suppressed prostaglandin synthesis as effectively as the parent drug.<sup>26-30</sup> All these NO-releasing NSAIDs have a nitrooxyalkyl group as the NO-releasing group. However, an important drawback to this design is the fact that production of NO from organic nitrate esters requires a three-electron reduction, and this metabolic activation decreases in efficiency on continued use of the drugs, contributing to

25

30

"nitrate tolerance".<sup>31</sup> In this regard, *O*<sup>2</sup>-unsubstituted *N*-diazene-1-ium-1,2-diols have the potential to release up to 2 equivalents of NO with half-lives that correlate well with their pharmacological durations of action. These observations suggest that *N*-diazene-1-ium-1,2-diols are minimally affected by metabolism, and are essentially different from currently available clinical vasodilators that require redox activation before NO is released.<sup>32</sup> *N*-diazene-1-ium-1,2-diols possess three attributes that make them especially attractive for designing drugs to treat a variety of disease states, namely structural diversity, dependable rates of NO release, and rich derivatization chemistry that facilitates targeting of NO to specific target organ and/or tissue sites.<sup>32</sup> As part of our ongoing research program to develop anti-inflammatory agents with a greater safety profile, Applicants now report the synthesis, *in vitro* COX-1/COX-2 inhibitory activity, *in vivo* anti-inflammatory activity, nitric oxide release data, and results from ulcerogenicity studies for a group of ester prodrugs of aspirin, ibuprofen and indomethacin that possess a diazene-1-ium-1,2-diolate as the NO-donor moiety.

#### SUMMARY OF THE INVENTION

The invention is intended to help protect chronic NSAID users such as arthritis and cardiovascular patients from potentially life-threatening gastrointestinal side effects without compromising anti-inflammatory activity. It provides a method of forming hybrid prodrugs comprising a non-steroidal anti-inflammatory drug (NSAID) linked by a methylene spacer on its carboxylic acid group to a diazene-1-ium-1,2-diolate moiety which on hydrolysis will release nitric oxide. It is intended to prevent or ameliorate gastrointestinal upset, bleeding or ulceration through the protective effect of nitric oxide in the tissues lining the gastrointestinal tract.

## DETAILED DESCRIPTION OF THE FIGURES

5 **Figure 1.** Chemical structures of acetyl salicylic acid (1) and some representative NO-NSAIDs (organic nitrates): NO-aspirin (2), NO-naproxen (3), NO-flurbiprofen (4) and NO-dichlofenac (5).

10 **Figure 2.** Ulcerogenicity assay data illustrating the extent of NSAID-induced gastric ulcers for NO-NSAIDs 11, 13 and 15, compared to that induced by the parent drugs aspirin, ibuprofen and indomethacin.

15 **Figure 3.** O<sup>2</sup>-Chloromethyl-1-(N,N-dimethylamino)diazene-1-ium-1,2-diolate (9) preparation procedure.

**Figure 4.** Synthesis of the target NO-NSAID ester prodrugs.

20 **Figure 5.** Theoretical metabolic activation (hydrolysis) of NO-NSAIDs (compound 13 shown as a representative example)

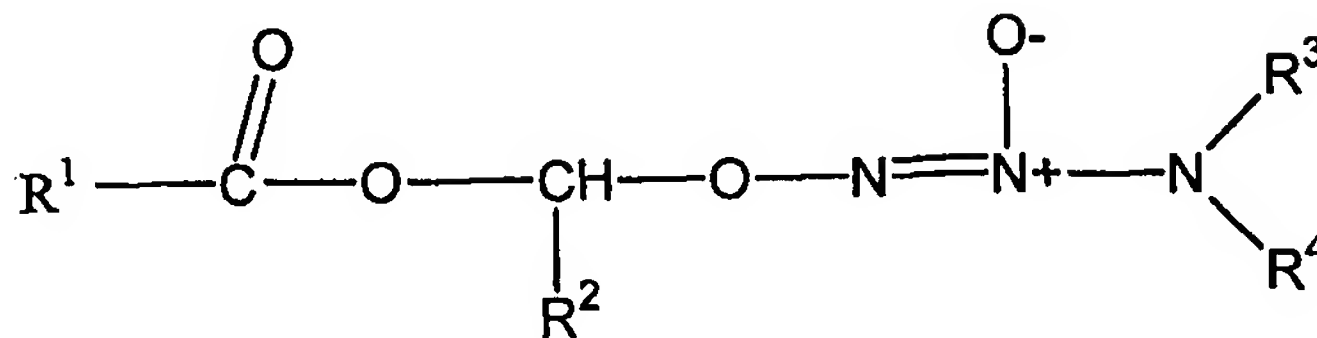
**Figure 6.** Structures of new NO-releasing non-steroidal anti-inflammatory drugs based on aspirin, ibuprofen and indomethacin (NO-NSAIDs)

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## DETAILED DESCRIPTION OF THE INVENTION

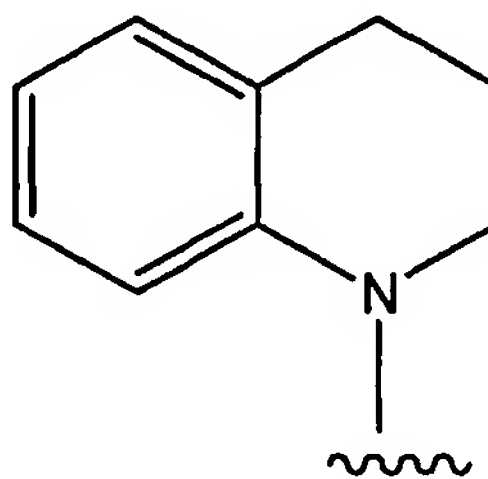
This invention provides a compound of the formula I:



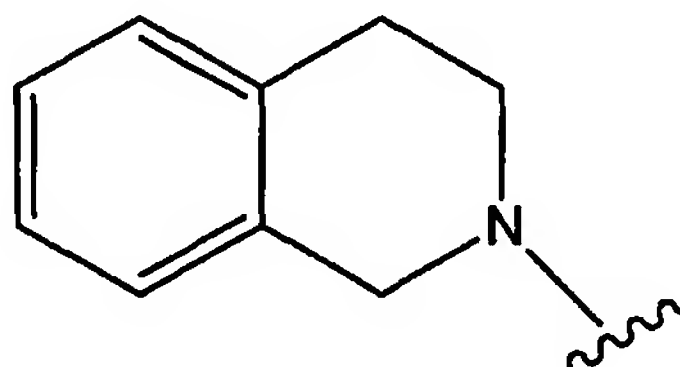
Structure I

wherein  $\text{R}^1$  is the uncarboxylated core of a non-steroidal anti-inflammatory drug,  $\text{R}^2$  is hydrogen, an unsubstituted or substituted  $\text{C}_{1-12}$  straight chain alkyl, an unsubstituted or substituted  $\text{C}_{3-12}$  branched chain alkyl, an unsubstituted or substituted  $\text{C}_{3-12}$  straight chain alkenyl, an unsubstituted or substituted  $\text{C}_{3-12}$  branched chain alkenyl, an unsubstituted or substituted  $\text{C}_{3-8}$  cycloalkyl, an unsubstituted or substituted alkoxy, nitrile, halo, an unsubstituted or substituted morpholino, amino, an unsubstituted or substituted benzyl, an unsubstituted or substituted phenyl, an unsubstituted or substituted  $\text{C}_{1-4}$  aryl alkyl, an unsubstituted or substituted heteroaryl, an unsubstituted or substituted arylamino, an unsubstituted or substituted dialkylamino, an unsubstituted or substituted diarylamino, carboxyalkylamino, carboxydialkylamino, an unsubstituted or substituted tolyl, xylyl, anisyl, mesityl, an unsubstituted or substituted acetoxy, carboxy, an unsubstituted or substituted carboxyethyl, an unsubstituted or substituted alkylcarbonyl, thiol, an unsubstituted or substituted alkylthiol, an unsubstituted or substituted alkyloxy, carboxyamido, an unsubstituted or substituted alkylcarboxyamido, an unsubstituted or substituted dialkylcarboxyamido, an unsubstituted or substituted phenoxy, an unsubstituted or

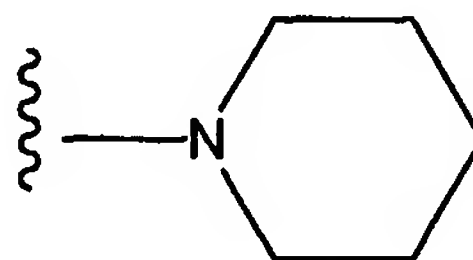
substituted benzyloxy, phenylcarbonyl, benzylcarbonyl, an  
unsubstituted or substituted nitrophenyl, trialkylsilyl or  
nitro;  $R^3$  and  $R^4$  are the same or different and are each  
preferentially one of an unsubstituted or substituted  $C_{1-12}$   
5 straight chain alkyl, an unsubstituted or substituted  $C_{3-12}$   
branched chain alkyl, an unsubstituted or substituted  $C_{3-12}$   
straight chain alkenyl, an unsubstituted or substituted  $C_{3-12}$   
branched chain alkenyl, an unsubstituted or substituted  $C_{3-8}$   
cycloalkyl, an unsubstituted or substituted morpholino, amino,  
10 an unsubstituted or substituted benzyl, an unsubstituted or  
substituted  $C_{1-4}$  aryl alkyl, , an unsubstituted or substituted  
carboxyethyl, or the  $-N(R^3, R^4)$  group is cyclized to form a  
1,2,3,4-tetrahydroquinolyl, i.e. Structure II:



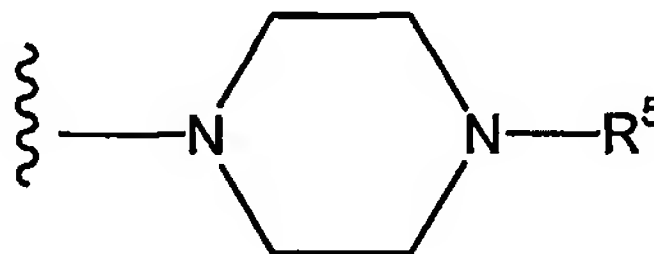
15 or structure III:



or piperidinyl, Structure IV:



or N-substituted-piperizinyll, Structure V:



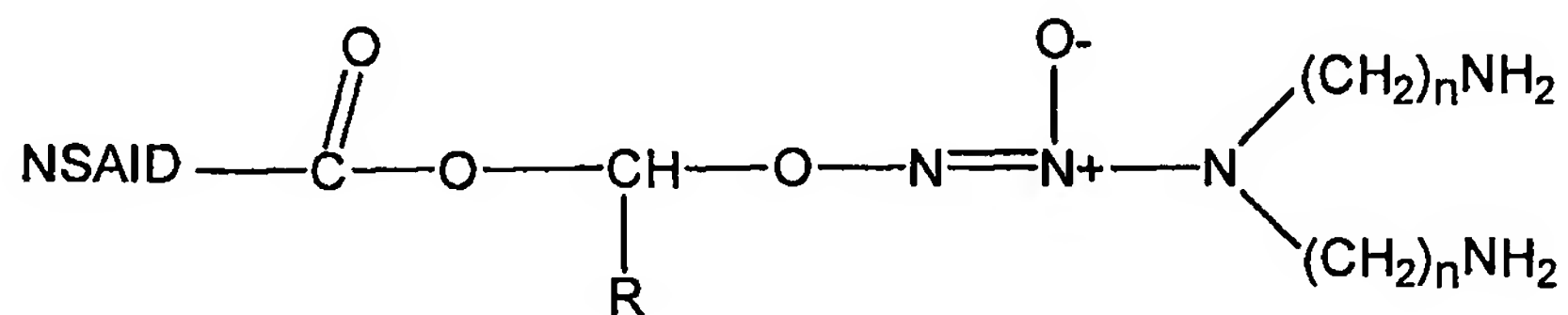
20



where  $R^5$  is an unsubstituted or substituted  $C_{1-12}$  straight chain alkyl, an unsubstituted or substituted  $C_{3-12}$  branched chain alkyl, an unsubstituted or substituted  $C_{3-12}$  straight chain alkenyl, an unsubstituted or substituted  $C_{3-12}$  branched chain alkenyl, an unsubstituted or substituted  $C_{3-8}$  cycloalkyl, an unsubstituted or substituted benzyl, an unsubstituted or substituted phenyl, an unsubstituted or substituted  $C_{1-4}$  aryl alkyl, an unsubstituted or substituted heteroaryl, an unsubstituted or substituted tolyl, xylyl, anisyl, mesityl, an unsubstituted or substituted carboxyethyl, an unsubstituted or substituted alkylcarbonyl, phenylcarbonyl, benzylcarbonyl, an unsubstituted or substituted nitrophenyl, or trialkylsilyl.

This invention also provides a compound of the formula I, wherein the non-steroidal anti-inflammatory drug carboxylic acid in  $R^1$  is acetylsalicylic acid, ibuprofen, naproxen, indomethacin, salicylic acid, diflunisal, salsalate, olsalazine, sulfasalazine, sulindac, etodolac, mefenamic acid, meclofenamic acid, tolmetin, ketorolac, diclofenac, fenoprofen, ketoprofen, oxaprozin, carprofen, flurbiprofen, nabumetone, any other related carboxylic acids with anti-inflammatory activity and their pharmaceutically suitable salts.

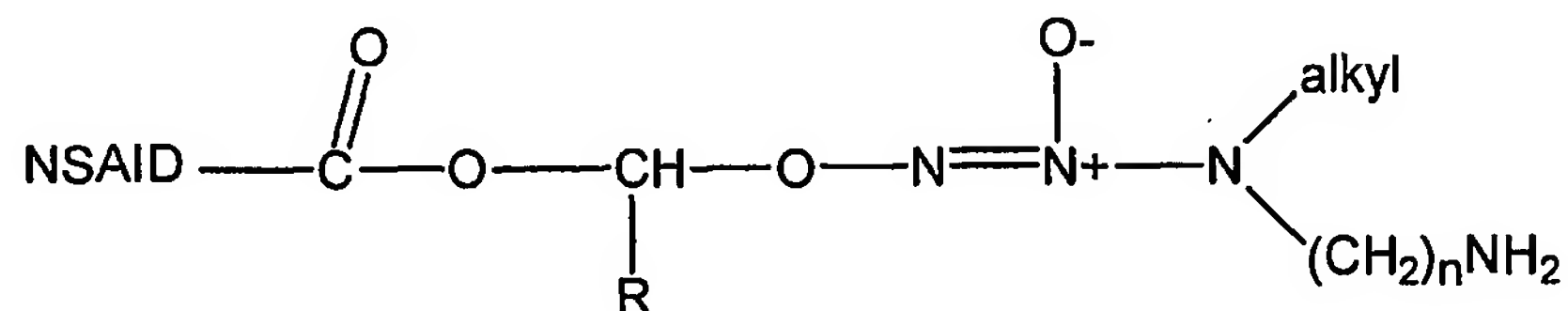
This invention provides a compound of the formula VII:



Structure VII

Wherein R is as in R<sup>2</sup> of Structure I, n=1-8. The structure includes pharmaceutically suitable alkali metal salts or hydrochloride salts of VII.

5 This invention provides a compound of Structure VIII:



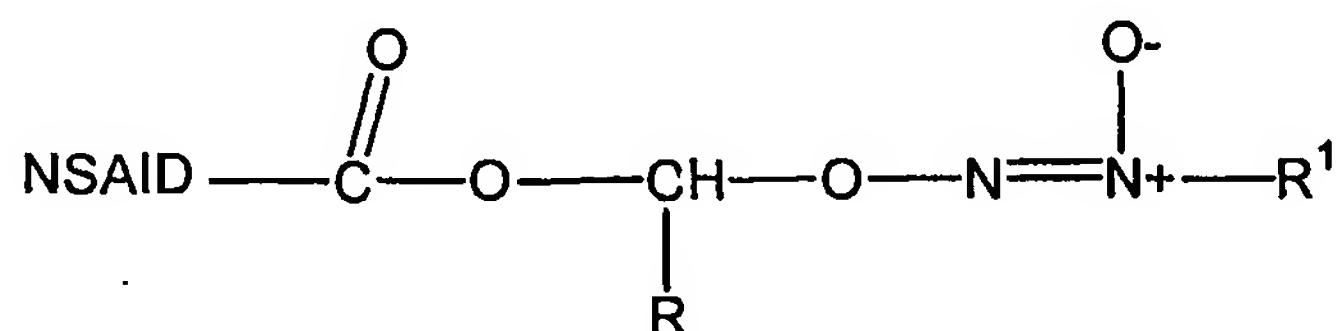
Structure VIII

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Wherein R is as in R<sup>2</sup> of Structure I, n=1-8. The structure includes pharmaceutically suitable alkali metal salts or hydrochloride salts of VIII.

15

This invention provides a compound of Structure IX:

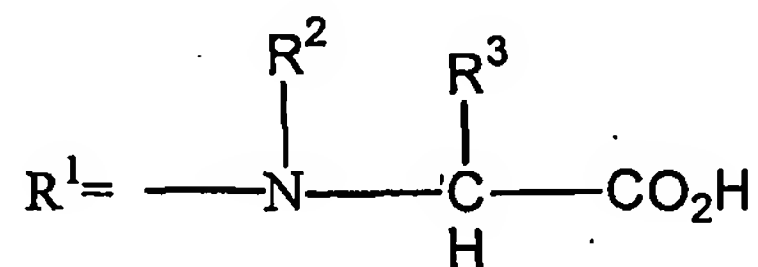


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Structure IX

Wherein R is as in R<sup>2</sup> of Structure I, R<sup>1</sup> is a N-substituted amino acid moiety.

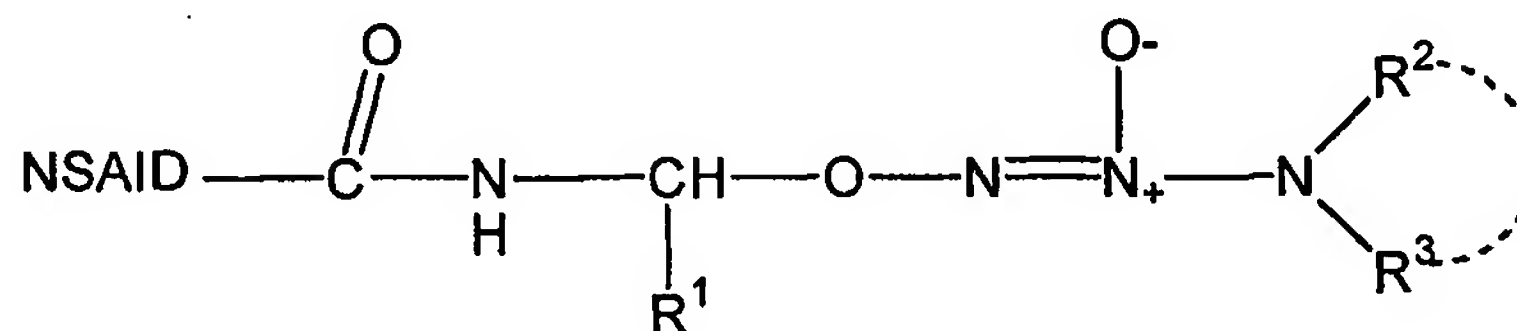
This invention provides a compound of Structure IX above,  
25 wherein R<sup>1</sup> the N-substituted amino acid moiety is:



And  $R^2$  is hydrogen, an unsubstituted or substituted  $C_{1-12}$  straight chain alkyl, an unsubstituted or substituted  $C_{3-12}$  branched chain alkyl, an unsubstituted or substituted  $C_{3-12}$  straight chain alkenyl, an unsubstituted or substituted  $C_{3-12}$  branched chain alkenyl, an unsubstituted or substituted  $C_{3-8}$  cycloalkyl, an unsubstituted or substituted benzyl, an unsubstituted or substituted phenyl, an unsubstituted or substituted  $C_{1-4}$  aryl alkyl, an unsubstituted or substituted heteroaryl, an unsubstituted or substituted tolyl, xylyl, anisyl, mesityl, an unsubstituted or substituted carboxyethyl, and  $R^3$  is hydrogen, an unsubstituted or substituted  $C_{1-12}$  straight chain alkyl, an unsubstituted or substituted  $C_{3-12}$  branched chain alkyl, an unsubstituted or substituted  $C_{3-12}$  straight chain alkenyl, an unsubstituted or substituted  $C_{3-12}$  branched chain alkenyl, an unsubstituted or substituted  $C_{3-8}$  cycloalkyl, an unsubstituted or substituted alkoxy, nitrile, halo, an unsubstituted or substituted morpholino, amino, an unsubstituted or substituted benzyl, an unsubstituted or substituted phenyl, an unsubstituted or substituted  $C_{1-4}$  aryl alkyl, an unsubstituted or substituted heteroaryl, an unsubstituted or substituted arylamino, an unsubstituted or substituted dialkylamino, an unsubstituted or substituted diarylamino, carboxyalkylamino, carboxydialkylamino, an unsubstituted or substituted tolyl, xylyl, anisyl, mesityl, an unsubstituted or substituted acetoxy, carboxy, an unsubstituted or substituted carboxyethyl, an unsubstituted or substituted alkylcarbonyl, an unsubstituted or substituted alkylthiol, an unsubstituted or substituted alkyloxy, carboxyamido, an unsubstituted or substituted alkylcarboxyamido, an unsubstituted or substituted dialkylcarboxyamido, an unsubstituted or substituted phenoxy, an unsubstituted or substituted benzyloxy, phenylcarbonyl, benzylcarbonyl, an unsubstituted or substituted nitrophenyl, trialkylsilyl or nitro. The simplest examples are N-

methylglycine, N-methylalanine, N-methylphenylalanine, N-methylserine, or any other N-alkyl amino acid.

This invention provides an amide bioisostere ester compound of structure X:



Structure X

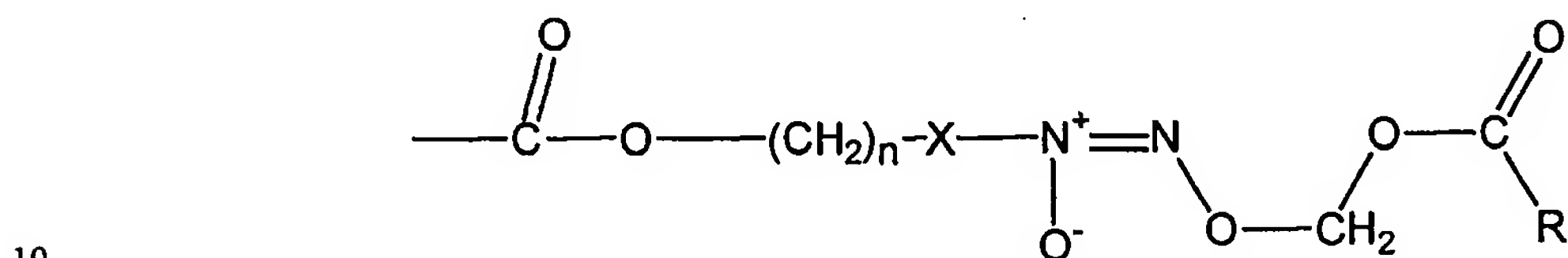
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Wherein  $R^1$  is hydrogen, an unsubstituted or substituted  $C_{1-12}$  straight chain alkyl, an unsubstituted or substituted  $C_{3-12}$  branched chain alkyl, an unsubstituted or substituted  $C_{3-12}$  straight chain alkenyl, an unsubstituted or substituted  $C_{3-12}$  branched chain alkenyl, an unsubstituted or substituted  $C_{3-8}$  cycloalkyl, an unsubstituted or substituted alkoxy, nitrile, halo, an unsubstituted or substituted morpholino, amino, an unsubstituted or substituted benzyl, an unsubstituted or substituted phenyl, an unsubstituted or substituted  $C_{1-4}$  aryl alkyl, an unsubstituted or substituted heteroaryl, an unsubstituted or substituted arylamino, an unsubstituted or substituted dialkylamino, an unsubstituted or substituted diarylamino, carboxyalkylamino, carboxydialkylamino, an unsubstituted or substituted tolyl, xylyl, anisyl, mesityl, an unsubstituted or substituted acetoxy, carboxy, an unsubstituted or substituted carboxyethyl, an unsubstituted or substituted alkylcarbonyl, thiol, an unsubstituted or substituted alkylthiol, an unsubstituted or substituted alkyloxy, carboxyamido, an unsubstituted or substituted alkylcarboxyamido, an unsubstituted or substituted dialkylcarboxyamido, an unsubstituted or substituted phenoxy,

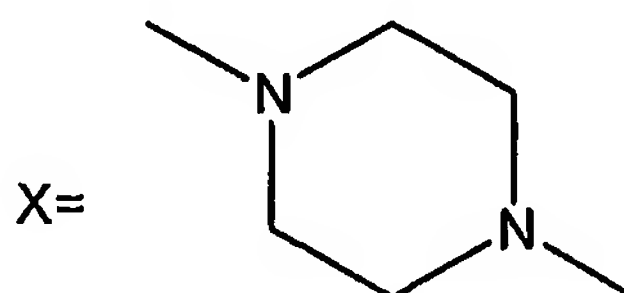
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an unsubstituted or substituted benzyloxy, phenylcarbonyl, benzylcarbonyl, an unsubstituted or substituted nitrophenyl, trialkylsilyl or nitro and the  $-N(R^2, R^3)$  group is cyclized to form a 1,2,3,4-tetrahydroquinolyl (Structure II above or structure III above), piperidinyl (Structure above) or N-substituted-piperizinyll (Structure V above).

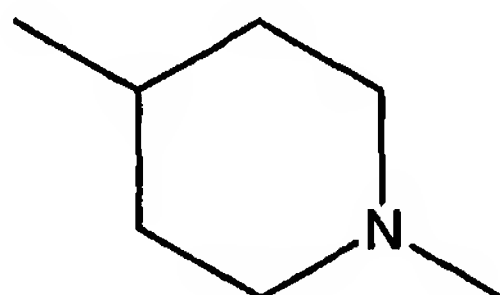
This invention provides A compound of structure XI:



Wherein X is a N-substituted piperizinyll



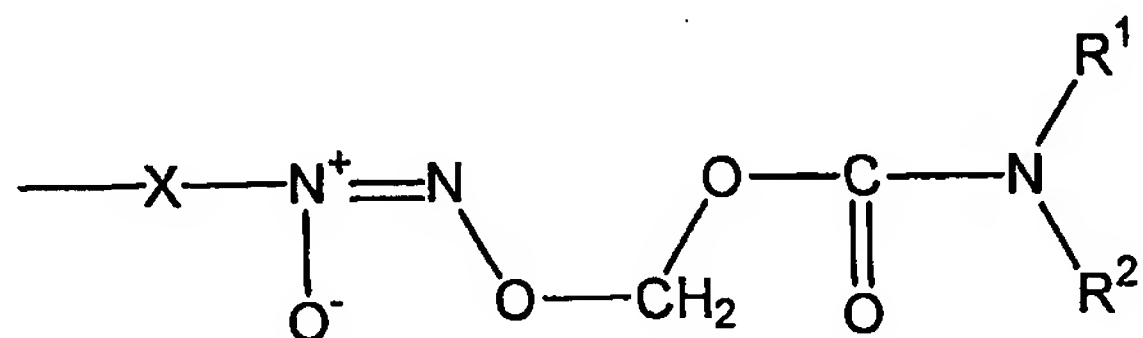
or N- and 4-substituted piperidinyl



20 or N-methyl moiety and R is an unsubstituted or substituted  $C_{1-12}$  straight chain alkyl, an unsubstituted or substituted  $C_{3-12}$  branched chain alkyl, an unsubstituted or substituted  $C_{3-12}$  straight chain alkenyl, an unsubstituted or substituted  $C_{3-12}$  branched chain alkenyl, an unsubstituted or substituted  $C_{3-8}$  cycloalkyl, an unsubstituted or substituted alkoxy, an  
25 unsubstituted or substituted morpholino, amino, an

unsubstituted or substituted benzyl, an unsubstituted or  
 substituted phenyl, an unsubstituted or substituted C<sub>1-4</sub> aryl  
 alkyl, an unsubstituted or substituted heteroaryl, an  
 unsubstituted or substituted arylamino, an unsubstituted or  
 5 substituted dialkylamino, an unsubstituted or substituted  
 diarylamino, carboxyalkylamino, carboxydialkylamino, an  
 unsubstituted or substituted tolyl, xylyl, anisyl, mesityl, an  
 unsubstituted or substituted acetoxy, carboxy, an  
 unsubstituted or substituted carboxyethyl, an unsubstituted or  
 10 substituted alkylcarbonyl, an unsubstituted or substituted  
 alkylthiol, an unsubstituted or substituted  
 alkyloxy, carboxyamido, an unsubstituted or substituted  
 alkylcarboxyamido, an unsubstituted or substituted  
 dialkylcarboxyamido, an unsubstituted or substituted phenoxy,  
 15 an unsubstituted or substituted benzyloxy, phenylcarbonyl,  
 benzylcarbonyl, an unsubstituted or substituted nitrophenyl,  
 trialkylsilyl or nitro.

This invention provides a carbamate compound of structure XII:



Structure XII

25

Wherein X is a N-substituted piperizinyll as in Structure XI,  
 a N- and 4-substituted piperidinyll as in Structure XI or N-  
 methylmoiety and R<sup>1</sup> and R<sup>2</sup> are each preferentially one of  
 hydrogen, an unsubstituted or substituted C<sub>1-12</sub> straight chain  
 30 alkyl, an unsubstituted or substituted C<sub>3-12</sub> branched chain  
 alkyl, an unsubstituted or substituted C<sub>3-12</sub> straight chain



alkenyl, an unsubstituted or substituted C<sub>3-12</sub> branched chain  
alkenyl, an unsubstituted or substituted C<sub>3-8</sub> cycloalkyl, an  
unsubstituted or substituted benzyl, an unsubstituted or  
substituted phenyl, an unsubstituted or substituted C<sub>1-4</sub> aryl  
5 alkyl, , an unsubstituted or substituted heteroaryl, an  
unsubstituted or substituted tolyl, xylyl, anisyl, mesityl, an  
unsubstituted or substituted carboxyethyl, an unsubstituted or  
substituted alkylcarbonyl, phenylcarbonyl, benzylcarbonyl, an  
unsubstituted or substituted nitrophenyl, or nitro or the -  
10 N(R<sup>2</sup>, R<sup>3</sup>) group is cyclized to form a 1,2,3,4-  
tetrahydroquinolyl (Structure II above or structure III  
above), piperidinyl (Structure IV above), or N-substituted-  
piperizinyl (Structure V above).

15 This invention provides a compound O<sup>2</sup>-  
(Acetylsalicyloyloxymethyl)-1-(pyrrolidin-1-yl)diazene-1-ium-  
1,2-diolate as shown in Figure 6.

This invention provides a compound O<sup>2</sup>-  
20 (Acetylsalicyloyloxymethyl)-1-(N,N-dimethylamino)diazene-1-ium-  
1,2-diolate as shown in Figure 6.

This invention provides a compound O<sup>2</sup>-[2-(4-  
(Isobutyl)phenyl)propanoyloxymethyl]-1-(pyrrolidin-1-yl)  
25 diazene-1-ium-1,2-diolate as shown in Figure 6.

This invention provides a compound O<sup>2</sup>-[2-(4-  
(Isobutyl)phenyl)propanoyloxymethyl]-1-(N,N-dimethylamino)  
diazene-1-ium-1,2-diolate as shown in Figure 6.

30 This invention provides a compound O<sup>2</sup>-[2-(1-(4-Chlorobenzoyl)-  
5-methoxy-2-methyl-1H-indol-3-yl)acetoxymethyl]-1-  
(pyrrolidin-1-yl)diazene-1-ium-1,2-diolate as shown in Figure  
6.

This invention provides a compound O<sup>2</sup>-[2-(1-(4-Chlorobenzoyl)-5-methoxy-2-methyl-1H-indol-3-yl)acetoxymethyl]-1-(dimethylamino)diazen-1-ium-1,2-diolate as shown in Figure 6.

5

This invention provides a composition comprising an effective amount of one of the compounds described herein in the same molar dose range as recommended for the NSAID from which it was derived.

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This invention provides a composition comprising an effective amount of one of the compounds described herein in various dose ranges capable of enhancing therapeutic outcome as recommended for the NSAID from which it was derived.

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This invention provides the use of any of the above-mentioned compounds to reduce gastrointestinal side effects of the parent non-steroidal anti-inflammatory drugs (NSAID). The side effects include but are not limited to dyspepsia, nausea and vomiting, abdominal pain, diarrhea, gastric or intestinal bleeding, and gastric and/or intestinal ulceration.

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This invention provides the use of any of the above-mentioned compounds for the indications recommended for the unsubstituted NSAID from which it is derived. For example the indication may be pain and inflammation, headache (e.g. ibuprofen), cardiovascular protection (e.g. acetylsalicylic acid), rheumatoid or osteoarthritis symptoms (e.g. naproxen, indomethacin), etc.

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This invention provides the use of any of the above-mentioned compounds in the same molar dose range as recommended for the NSAID from which it was derived.

This invention provides the use of any of the above-mentioned compounds described in various dose ranges to achieve better therapeutic outcome as recommended for the NSAID from which it was derived.

5

#### EXEMPLIFICATION

The invention being generally described, will be more readily understood by reference to the following examples which are included merely for purposes of illustration of certain aspects and embodiments of the present invention, and are not intended to limit the invention.

A group of new NO-releasing non-steroidal anti-inflammatory drugs (NO-NSAIDs), derived from aspirin ( $O^2$ -(Acetylsalicyloyloxymethyl)-1-(pyrrolidin-1-yl)diazen-1-ium-1,2-diolate, 11;  $O^2$ -(Acetylsalicyloyloxymethyl)-1-(N,N-dimethylamino)diazen-1-ium-1,2-diolate, 12), ibuprofen ( $O^2$ -[2-(4-(Isobutyl)phenyl)propanoyloxymethyl]-1-(pyrrolidin-1-yl)diazen-1-ium-1,2-diolate, 13;  $O^2$ -[2-(4-(Isobutyl)phenyl)propanoyloxymethyl]-1-(N,N-dimethylamino)diazen-1-ium-1,2-diolate, 14) and indomethacin ( $O^2$ -[2-(1-(4-Chlorobenzoyl)-5-methoxy-2-methyl-1H-indol-3-yl)acetoxymethyl]-1-(pyrrolidin-1-yl)diazen-1-ium-1,2-diolate, 15;  $O^2$ -[2-(1-(4-Chlorobenzoyl)-5-methoxy-2-methyl-1H-indol-3-yl)acetoxymethyl]-1-(dimethyl amino)diazen-1-ium-1,2-diolate, 16) possessing a 1-(pyrrolidin-1-yl)diazen-1-ium-1,2-diolate, or 1-(N,N-dimethylamino)diazen-1-ium-1,2-diolate moiety were synthesized.

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Chemistry:  $O^2$ -Chloromethyl-1-(N,N-dimethylamino)diazen-1-ium-1,2-diolate (9) was prepared according to a modified procedure reported by Tang et al,<sup>33</sup> as illustrated in Figure 3. Thus, reaction of dimethylamine (6) with nitric oxide gas (40 psi)

at room temperature in the presence of sodium methoxide, afforded O<sup>2</sup>-sodium 1-(N,N-dimethylamino)diazene-1-ium-1,2-diolate (7) in 90% yield. The sodium salt was alkylated with chloromethyl methyl sulfide to afford O<sup>2</sup>-(methylthiomethyl)-1-(N,N-dimethylamino)diazene-1-ium-1,2-diolate (8), which was subsequently reacted with sulfuryl chloride in dichloromethane for 4 h to afford the O<sup>2</sup>-chloromethyl-protected diazeniumdiolate 9 in quantitative yield. The target NO-NSAID ester prodrugs 11-16 were synthesized in moderate-to-good yields (40-81%) by condensation of the sodium salt of acetylsalicylic acid, ibuprofen or indomethacin, with O<sup>2</sup>-chloromethyl intermediates 9 or 10 using the polar aprotic solvent HMPA (Figure 4).

In vitro COX enzyme inhibition studies, showed that none of these compounds inhibited either the COX-1 or COX-2 isozyme at the highest test compound concentration used (100  $\mu$ M). See Table 1 below.

**Table 1.** In Vitro COX-1/COX-2 Enzyme Inhibition, and In Vivo Antiinflammatory Activity Data for NO-NSAIDs 11-16.

Compd.	COX-1 IC <sub>50</sub> ( $\mu$ M) <sup>a</sup>	COX-2 IC <sub>50</sub> ( $\mu$ M) <sup>a</sup>	COX-2 S.I. <sup>b</sup>	AI activity <sup>c</sup> ID <sub>50</sub> (mg/kg)
11	> 100	> 100	-	181.8
12	> 100	> 100	-	151.2
13	> 100	> 100	-	66.8
14	> 100	> 100	-	62.3
15	> 100	> 100	-	10.7
16	> 100	> 100	-	5.9
Aspirin	0.3	2.4	0.14	128.7

Ibuprofen	2.9	1.1	2.63	67.4
Indomethacin	0.1	5.7	0.01	4.2

- 5 a The in vitro test compound concentration required to produce 50% inhibition of COX-1 or COX-2. The result ( $IC_{50}$ ,  $\mu M$ ) is the mean of two determinations acquired using an ovine COX-1/COX-2 assay kit (Catalog No. 560101, Cayman Chemicals Inc., Ann Arbor, MI, USA) and the deviation from the mean is <10% of the mean value.
- b Selectivity index (SI) =  $COX-1\ IC_{50}/COX-2\ IC_{50}$ .
- 10 c Inhibitory activity in a carrageenan-induced rat paw edema assay. The results are expressed as the  $ID_{50}$  value (mg/kg) at 3 h after oral administration of the test compound.

Thus, attachment of an ester group (the NO-releasing diazeniumdiolate moiety) to the parent NSAID completely abolished the in vitro enzyme inhibitory activity of aspirin, 15 ibuprofen and indomethacin. However, when administered orally to rats, the carrageenan-induced rat paw edema assay (Table 1) provided similar  $ID_{50}$  values to those obtained for the reference drugs. The ibuprofen NO-NSAIDs 13 and 14 showed equipotent anti-inflammatory activities ( $ID_{50}$  = 66.8 and 62.3 20 mg/kg respectively) compared to the reference drug ibuprofen ( $ID_{50}$  = 67.4 mg/kg). Similar results were obtained for the NO-aspirins 11 ( $ID_{50}$  = 181.8 mg/kg) and 12 ( $ID_{50}$  = 151.2 mg/kg), and the NO-indomethacin 16 ( $ID_{50}$  = 5.9 mg/kg), which were 1.1- 25 = 128.7 mg/kg) and indomethacin ( $ID_{50}$  = 4.2 mg/kg). In comparison, the NO-indomethacin 15 ( $ID_{50}$  = 10.7 mg/kg) was about 2.5 fold-less potent than indomethacin. Compounds containing a 1-(pyrrolidin-1-yl)diazen-1-ium-1,2-diolate (11, 13 and 15) moiety were less active than those compounds having 30 a 1-(N,N-dimethylamino)diazen-1-ium-1,2-diolate moiety (12, 14 and 16). It has been reported that aspirin acetylates the Ser530 residue in the COX-1 active site.<sup>14</sup> The observations that both NO-aspirins (11 and 12) were inactive in vitro inhibitors of COX-1 and COX-2 ( $IC_{50}$  > 100  $\mu M$ ), and that they

showed significant anti-inflammatory activities in vivo, strongly suggests that 11 and 12 act as classical prodrugs, which require a metabolic activation reaction (esterase-mediated ester cleavage) to be active. One type of chemical modification used to control the rate of nitric oxide release from diazen-1-ium-1,2-diolates is the attachment of alkyl substituents to the O<sup>2</sup>-position.<sup>34</sup> O<sup>2</sup>-substituted-diazen-1-ium-1,2-diolates are stable compounds that hydrolyze slowly even in acidic solution.<sup>35</sup> Consistent with these observations, when compounds 11-16 were incubated in phosphate buffer solution (PBS) at pH 7.4, the percentage of NO released varied from 14.3 to 16.1 % which is indicative of slow NO release. In contrast to recently reported O<sup>2</sup>-acetoxymethyl-1-(pyrrolidin-1-yl or N,N-diethylamino)diazen-1-ium-1,2-diolates,<sup>36</sup> which are stable prodrugs in neutral aqueous media but which released about 1.8 equivalents of NO (> 90% release) per mol of drug upon metabolism by porcine liver esterase (PLE), the ester prodrugs 11-16 are hydrolyzed much less extensively (16.3 to 19.2% NO release). However, the effect of non-specific esterases present in guinea pig serum on the NO release properties of compounds 11-16 was substantially higher (81.6-93.6% range) than that observed (16.3-19.2% range) upon incubation with PLE (see Table 2).

Table 2. Nitric Oxide Release Data for NO-NSAIDs 11-16.

Compd	% of Nitric oxide released <sup>a</sup>		
	PBS (pH 7.4) <sup>b</sup>	PLE <sup>c</sup>	GP-Serum <sup>d</sup>
11	14.8 ± 0.1	18.5 ± 0.1	88.9 ± 0.2
12	15.4 ± 0.1	19.1 ± 0.1	81.6 ± 0.1
13	14.9 ± 0.1	16.3 ± 0.1	89.2 ± 0.1
14	16.1 ± 0.1	17.3 ± 0.1	93.6 ± 0.1
15	15.1 ± 0.1	16.3 ± 0.1	89.1 ± 0.1
16	14.3 ± 0.1	16.9 ± 0.1	86.3 ± 0.1
7	95.2 ± 0.1	-	-



O<sup>2</sup>-sodium 1- 94.0 + 0.1 - -  
 (pyrrolidin-1-yl)  
 diazen-1-ium-1,2-  
 diolate

- 
- a Percent of nitric oxide released ( $\pm$  SEM, n = 3) relative to a theoretical maximum release of 2 mol of NO/mol of test compound.
- b Incubated in phosphate buffer solution (PBS, pH 7.4) at 37°C for 1.5 h.
- c Incubated in the presence of 2 equivalents of pig liver esterase (based on a ratio of 1 mol of test compound / 2 mol of esterase) in phosphate buffer solution (pH 7.4) at 37°C for 1.5 h.
- d Test compound ( $2.0 \times 10^{-4}$  mmol) incubated with guinea pig serum (260  $\mu$ L) in phosphate buffer solution (pH 7.4) at 37°C for 1.5 h.
- These data indicate the non-specific serum esterases present in guinea pig serum cleave these NO-NSAIDs more effectively than PLE. Although conventional NO donors can protect the stomach against NSAID-induced gastric damage, they do not do so as effectively as NSAIDs (including aspirin) that are chemically linked to an NO-releasing moiety.<sup>37</sup> A plausible mechanism for the hydrolysis of these NO-NSAID ester prodrugs 11-16 is presented in Figure 5. The NO-NSAID ester prodrugs 11-16 were designed with a one-carbon methylene spacer between the carboxy group and the diazen-1-ium-1,2-diolate O<sup>2</sup>-atom, such that the O<sup>2</sup>-(hydroxymethyl)diazen-1-ium-1,2-diolate compound formed after ester cleavage would spontaneously eliminate formaldehyde to produce the free NONOate compound that can subsequently fragment to release two molecules of NO.

One of the common side effects of NSAID therapy is gastrointestinal irritation and bleeding. It was therefore essential to evaluate the prodrugs 11-16 ulcerogenicity in comparison to that induced by the three parent drugs. The severity of gastric damage was expressed as an ulcer index (Table 3).

**Table 3.** Gastric ulcer index produced by an acute administration of the test compounds 11-16 and the reference drugs aspirin, ibuprofen and indomethacin.

Compd.	Ulcer index <sup>a</sup>
aspirin	57.4 ± 3.1 <sup>b</sup>
ibuprofen	45.8 ± 2.9 <sup>b</sup>
indomethacin	34.4 ± 4.2 <sup>c</sup>
11	0 <sup>d</sup>
12	0 <sup>d</sup>
13	0 <sup>e</sup>
14	0 <sup>e</sup>
15	0.7 ± 0.11 <sup>f</sup>
16	3.0 ± 0.3 <sup>f</sup>
control group	0 <sup>g</sup>

5 a The average overall length (in mm) of individual ulcers in each stomach ± SEM, n = 4, at 6 h after oral administration of the test compound.

b 250 mg/kg dose.

c 30 mg/kg dose.

d Equivalent amount to 250 mg of aspirin/kg.

10 e Equivalent amount to 250 mg of ibuprofen/kg.

f Equivalent amount to 30 mg of indomethacin/kg.

g 1.0% methylcellulose solution.

There was a remarkable difference between the ulcer index values for the NO-NSAIDs (UI = 0-3.0), and the reference drugs aspirin (UI = 57.4, 250 mg/kg po dose), ibuprofen (UI = 45.7, 250 mg/kg po dose) and indomethacin (34.4, 30 mg/kg po dose). This UI data suggests a much more safer pharmacological profile for hybrid NO-NSAIDs containing either a 1-(pyrrolidin-1-yl or N,N-dimethylamino)diazen-1-ium-1,2-diolate groups, relative to the parent drugs. No evidence of gastric ulcerogenicity (UI = 0) was observed (Figure 2) for either the NO-aspirin (11, 12) and NO-ibuprofen (13, 14) ester prodrugs. The NO-indomethacin compounds (15, 16) caused minimal ulcerogenicity (UI = 0.7-3.0 range).

## Conclusions

Hybrid NO-NSAID ester prodrugs possessing a 1-(pyrrolidin-1-yl)diazen-1-ium-1,2-diolate (11, 13, 15) or 1-(N,N-dimethylamino)diazen-1-ium-1,2-diolate (12, 14, 16), moiety  
5 attached via a one-carbon methylene spacer to the carboxylic acid group of traditional NSAIDs constitutes a useful concept for the rational design of anti-inflammatory drugs with reduced gastric side effects (ulcerogenicity). Virtually every NSAID having a free carboxylic acid is suitable for  
10 application of this methodology. In vivo activation (hydrolysis) of these NO-NSAIDs by plasma esterases, rather than liver esterases, would be expected to improve the NO release profile compared to that observed for organic nitrates which require a more metabolically demanding three-electron  
15 reduction for the release of NO, or a thiol cofactor such as L-cysteine or glutathione required for the release of NO from furoxans. Hybrid NO-aspirins having a diazen-1-ium-1,2-diolate moiety could be a useful alternative to the use of aspirin as an antithrombotic agent (inhibition of platelet aggregation)  
20 in the long-term prophylactic prevention of stroke and myocardial infarction.

General. Melting points were recorded with a Thomas-Hoover capillary apparatus and are uncorrected. <sup>1</sup>H NMR spectra were  
25 acquired using a Bruker AM-300 spectrometer (300 MHz). Infrared spectra were recorded using a Nicolet IR-500 Series II spectrometer. Silica gel column chromatography was carried out using Merck 7734 (60-200 mesh) silica gel.

30 Microanalyses were within  $\pm 0.4\%$  of theoretical values for all elements listed. See Table 4 below.

## Table 4.

### Microanalytical Data

Compound	Empirical	Calculated			Found		
	Formula	C	H	N	C	H	N
11	C <sub>14</sub> H <sub>17</sub> N <sub>3</sub> O <sub>6</sub>	52.01	5.30	13.00	51.99	5.28	12.90
12	C <sub>12</sub> H <sub>15</sub> N <sub>3</sub> O <sub>6</sub>	48.48	5.09	14.14	48.78	4.97	14.01
13	C <sub>18</sub> H <sub>27</sub> N <sub>3</sub> O <sub>4</sub>	61.87	7.79	12.03	61.83	7.79	12.03
14	C <sub>16</sub> H <sub>25</sub> N <sub>3</sub> O <sub>4</sub>	59.42	7.79	12.99	59.41	7.80	12.89
15	C <sub>24</sub> H <sub>25</sub> ClN <sub>4</sub> O <sub>6</sub>	57.54	5.03	11.18	57.53	5.03	11.22
16	C <sub>22</sub> H <sub>23</sub> ClN <sub>4</sub> O <sub>6</sub>	55.64	4.88	11.80	55.63	4.89	11.79

Acetyl salicylic acid (aspirin), racemic ibuprofen and indomethacin were purchased from the Sigma Chemical Co. O<sup>2</sup>-(chloromethyl)diazen-1-ium-1,2-diolate (10) was prepared according to a literature procedure<sup>33</sup> except that the reaction of O<sup>2</sup>-sodium 1-(pyrrolidin-1-yl)diazen-1-ium-1,2-diolate with chloromethyl methyl sulfide was carried out in HMPA at 25 °C for 48 h. Nitric oxide gas was purchased from BOC Scientific (Burlington, ON). All other chemicals were purchased from the Aldrich Chemical Co. (Milwaukee, WI). The in vivo anti-inflammatory and ulcer index assays were carried out using protocols approved by the Health Sciences Animal Welfare Committee at the University of Alberta.

O<sup>2</sup>-Sodium 1-(N,N-dimethylamino)diazen-1-ium-1,2-diolate (7). Dimethylamine (6, 4.5 g, 0.1 mol) was added to a solution of sodium methoxide (0.1 mol, 24 mL of a 25% w/v solution in methanol) and diethyl ether (300 mL) with stirring at 25 °C. This mixture was flushed with dry nitrogen for five minutes and then the reaction was allowed to proceed under an atmosphere of nitric oxide (40 psi internal pressure) with stirring at 25 °C for 19 h. The product, which precipitated as a fine white powder, was isolated by filtration and then suspended in diethyl ether (100 mL) upon stirring for 15 min. The suspension was filtered, the solid collected was dried at

25 °C under reduced pressure until a constant weight was achieved after about 2 h to afford 7 as a fine white powder (11.5 g, 90 %); mp 258-260 °C (dec.); <sup>1</sup>H NMR (DMSO-d<sub>6</sub>) δ 2.97 [s, 6H, N(CH<sub>3</sub>)<sub>2</sub>]. Product 7 was used immediately after drying without further purification for the preparation of compound 8.

O<sup>2</sup>-(Methylthiomethyl)-1-(N,N-dimethylamino)diazen-1-ium-1,2-diolate (8). The sodium diazeniumdiolate 7 (7 g, 54.6 mmol) was added to a suspension of potassium carbonate (1.5 g, 11 mmol) and HMPA (80 mL) at 4 °C and this mixture was stirred for 30 min. Chloromethyl methyl sulfide (6.3 g, 65.6 mmol) was added drop wise, and the reaction was allowed to proceed at 25 °C for 72 h with stirring. Ethyl acetate (200 mL) was added to quench the reaction, the solids were filtered off and the organic phase was washed with water (5 x 80 mL), dried (Na<sub>2</sub>SO<sub>4</sub>), and solvent was removed in vacuo to give a liquid residue which was purified by silica gel column chromatography using EtOAc-hexane (1:4, v/v) as eluent. Compound 8 (1.97 g, 21%) was obtained as a pale yellow liquid; <sup>1</sup>H NMR (CDCl<sub>3</sub>) δ 2.24 (s, 3H, SCH<sub>3</sub>), 3.01 [s, 6H, N(CH<sub>3</sub>)<sub>2</sub>], 5.21 (s, 2H, OCH<sub>2</sub>S). Compound 8 was used immediately for the subsequent preparation of the O<sup>2</sup>-chloromethyl derivative 9.

O<sup>2</sup>-(Chloromethyl)-1-(N,N-dimethylamino)diazen-1-ium-1,2-diolate (9). A solution of compound 8 (1.8 g, 11.4 mmol) in dichloromethane (20 mL) was cooled to 4 °C, sulfuryl chloride (2.3 g, 17.1 mmol, 17 mL of a 1.0 M solution in dichloromethane) was added drop wise, the ice bath was removed and the reaction mixture was stirred at 25 °C for 3 h. The brown solid suspended in the reaction media was removed by filtration, and the solvent was evaporated to furnish 9 (1.7 g, quantitative yield); <sup>1</sup>H NMR (CDCl<sub>3</sub>) δ 3.01 [s, 6H, N(CH<sub>3</sub>)<sub>2</sub>],

5.76 (s, 2H, ClCH<sub>2</sub>O). Compound 9 was used without further purification for the synthesis of products 12, 14 and 16.

General Method for the Preparation of NO-NSAIDs (11-16).  
5 Sodium carboxylates of the respective NSAID (aspirin, ibuprofen or indomethacin) were prepared in situ by stirring each acid (5 mmol) in a suspension of sodium carbonate (0.53 g, 5 mmol) and HMPA (7 mL) for 19 h at 25 °C. A solution of a  
10 O<sup>2</sup>-(chloromethyl)diazen-1-ium-1,2-diolate 9 or 10 (5 mmol) in HMPA (3 mL) was then added, and the reaction was allowed to proceed for 24 h at 25 °C. Ethyl acetate (60 mL) was added, the mixture was washed with water (5 x 30 mL), the organic phase was dried (Na<sub>2</sub>SO<sub>4</sub>), and the solvent was removed in vacuo. The residue obtained was purified by silica gel column  
15 chromatography using CHCl<sub>3</sub>-EtOAc-hexane (35:15:50, v/v/v) as eluent for compounds 11, 12, 15, and 16; EtOAc-hexane (1:4, v/v) for compound 13; and hexane-ether (3:1, v/v) for compound 14. Physical and spectral data for 11-16 are listed below.

20 O<sup>2</sup>-(Acetylsalicyloyloxymethyl)-1-(pyrrolidin-1-yl)diazen-1-ium-1,2-diolate (11). 46 % yield; white crystals; mp 110-112 °C; IR (CHCl<sub>3</sub>) 3019 (C-H aromatic), 2992 (C-H aliphatic), 1770 (CO<sub>2</sub>), 1736 (CO<sub>2</sub>), 1259, 1199 (N=N-O) cm<sup>-1</sup>; <sup>1</sup>H NMR (CDCl<sub>3</sub>) δ 1.95 (quintet, J = 6.9 Hz, 4H, pyrrolidinyl H-3, H-4), 2.34  
25 (s, 3H, COCH<sub>3</sub>), 3.57 (t, J = 6.9 Hz, 4H, pyrrolidinyl H-2, H-5), 5.97 (s, 2H, OCH<sub>2</sub>O), 7.12 (d, J = 8.1 Hz, phenyl H-3), 7.34 (t, J = 8.1 Hz, phenyl H-5), 7.60 (td, J = 8.1, 1.5 Hz, phenyl H-4), 8.08 (dd, J = 8.1, 1.5 Hz, phenyl H-6). Anal. (C<sub>14</sub>H<sub>17</sub>N<sub>3</sub>O<sub>6</sub>) C, H, N.

30 O<sup>2</sup>-(Acetylsalicyloyloxymethyl)-1-(N,N-dimethylamino)diazen-1-ium-1,2-diolate (12). 40 % yield; white crystals; mp 88-89 °C; IR (KBr) 3019 (C-H aromatic), 2979 (C-H aliphatic), 1756 (CO<sub>2</sub>),



1609 (CO<sub>2</sub>), 1219, 1184 (N=N-O) cm<sup>-1</sup>; <sup>1</sup>H NMR (CDCl<sub>3</sub>) δ 2.34 (s, 3H, COCH<sub>3</sub>), 3.07 [s, 6H, N(CH<sub>3</sub>)<sub>2</sub>], 6.02 (s, 2H, OCH<sub>2</sub>O), 7.12 (d, J = 8.1 Hz, phenyl H-3), 7.34 (t, J = 8.1 Hz, phenyl H-5), 7.60 (td, J = 8.1, 1.5 Hz, phenyl H-4), 8.07 (dd, J = 8.1, 1.5 Hz, phenyl H-6). Anal. (C<sub>12</sub>H<sub>15</sub>N<sub>3</sub>O<sub>6</sub>) C, H, N.

O<sup>2</sup>-[2-(4-(Isobutyl)phenyl)propanoyloxymethyl]-1-(pyrrolidin-1-yl)diazen-1-ium-1,2-diolate (13). 58 % yield; yellow oil; IR (KBr) 2985 (C-H aromatic), 2864 (C-H aliphatic), 1750 (CO<sub>2</sub>), 1286, 1129 (N=N-O) cm<sup>-1</sup>; <sup>1</sup>H NMR (CDCl<sub>3</sub>) δ 0.89 [d, J = 6.6 Hz, 6H, CH(CH<sub>3</sub>)<sub>2</sub>], 1.50 (d, J = 7.2 Hz, 3H, PhCHCH<sub>3</sub>), 1.79-1.89 [m, 1H, CH(CH<sub>3</sub>)<sub>2</sub>], 1.91-1.94 (m, 4H, pyrrolidiny H-3, H-4), 2.43 (d, J = 7.2 Hz, 2H, PhCH<sub>2</sub>CH), 3.45-3.50 (m, 4H, pyrrolidiny H-2, H-5), 3.73 (q, J = 7.2 Hz, 1H, PhCHCH<sub>3</sub>), 5.71 (d, J = 7.2 Hz, 1H, OCH'HO), 5.77 (d, J = 7.2 Hz, 1H, OCH'HO), 7.07 (d, J = 7.8 Hz, 2H, phenyl H-3, H-5), 7.19 (d, J = 7.8 Hz, 2H, phenyl H-2, H6). Anal. (C<sub>18</sub>H<sub>27</sub>N<sub>3</sub>O<sub>4</sub>) C, H, N.

O<sup>2</sup>-[2-(4-(Isobutyl)phenyl)propanoyloxymethyl]-1-(N,N-dimethylamino)diazen-1-ium-1,2-diolate (14). 81 % yield; yellow oil; IR (KBr) 2959 (C-H aromatic), 2871 (C-H aliphatic), 1763 (CO<sub>2</sub>), 1279, 1138 (N=N-O) cm<sup>-1</sup>; <sup>1</sup>H NMR (CDCl<sub>3</sub>) δ 0.89 [d, J = 6.9 Hz, 6H, CH(CH<sub>3</sub>)<sub>2</sub>], 1.50 (d, J = 6.9 Hz, 3H, PhCHCH<sub>3</sub>), 1.83 [septet, J = 6.9 Hz, 1H, CH(CH<sub>3</sub>)<sub>2</sub>], 2.43 (d, J = 6.9 Hz, 2H, PhCH<sub>2</sub>CH), 2.97 [s, 6H, N(CH<sub>3</sub>)<sub>2</sub>], 3.74 (q, J = 6.9 Hz, 1H, PhCHCH<sub>3</sub>), 5.74 (d, J = 7.2 Hz, 1H, OCH'HO), 5.79 (d, J = 7.2 Hz, 1H, OCH'HO), 7.08 (d, J = 7.8 Hz, 2H, phenyl H-3, H-5), 7.19 (d, J = 7.8 Hz, 2H, phenyl H-2, H6). Anal. (C<sub>16</sub>H<sub>25</sub>N<sub>3</sub>O<sub>4</sub>) C, H, N.

O<sup>2</sup>-[2-(1-(4-Chlorobenzoyl)-5-methoxy-2-methyl-1H-indol-3-yl)acetoxymethyl]-1-(pyrrolidin-1-yl)diazen-1-ium-1,2-diolate (15). 51 % yield; yellow oil; IR (KBr) 3019 (C-H aromatic),

2979, 2885 (C-H aliphatic), 1756 (CON), 1689 (CO<sub>2</sub>), 1293, 1165 (N=N-O) cm<sup>-1</sup>; <sup>1</sup>H NMR (CDCl<sub>3</sub>) δ 1.88 (quintet, J = 6.9 Hz, 4H, pyrrolidinyl H-3, H-4), 2.36 (s, 3H, CH<sub>3</sub>), 3.40 (t, J = 6.9 Hz, 4H, pyrrolidinyl H-2, H-5), 3.71 (s, 2H, CH<sub>2</sub>CO<sub>2</sub>), 3.83 (s, 3H, OCH<sub>3</sub>), 5.77 (s, 2H, OCH<sub>2</sub>O), 6.66 (dd, J = 9, 2.4 Hz, 1H, indolyl H-6), 6.90 (d, J = 9 Hz, 1H, indolyl H-7), 6.94 (d, J = 2.4 Hz, indolyl H-4), 7.47 (d, J = 8.7 Hz, 2H, benzoyl H-3, H-5), 7.65 (d, J = 8.7 Hz, 2H, benzoyl H-2, H-6). Anal. (C<sub>24</sub>H<sub>25</sub>ClN<sub>4</sub>O<sub>6</sub>) C, H, N.

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O<sup>2</sup>-[2-(1-(4-Chlorobenzoyl)-5-methoxy-2-methyl-1H-indol-3-yl)acetoxymethyl]-1-(dimethyl amino)diazene-1-ium-1,2-diolate (16). 69 % yield; yellow oil; IR (KBr) 2979, 2925 (C-H aliphatic), 1763 (CON), 1689 (CO<sub>2</sub>), 1333, 1064 (N=N-O) cm<sup>-1</sup>; <sup>1</sup>H NMR (CDCl<sub>3</sub>) δ 2.35 (s, 3H, CH<sub>3</sub>), 2.94 [s, 6H, N(CH<sub>3</sub>)<sub>2</sub>], 3.71 (s, 2H, CH<sub>2</sub>CO<sub>2</sub>), 3.81 (s, 3H, OCH<sub>3</sub>), 5.80 (s, 2H, OCH<sub>2</sub>O), 6.66 (dd, J = 8.7, 2.4 Hz, 1H, indolyl H-6), 6.88 (d, J = 8.7 Hz, 1H, indolyl H-7), 6.93 (d, J = 2.4 Hz, 1H, indolyl H-4), 7.46 (d, J = 8.4 Hz, 2H, benzoyl H-3, H-5), 7.64 (d, J = 8.4, 2H, benzoyl H-2, H-6). Anal. (C<sub>22</sub>H<sub>23</sub>ClN<sub>4</sub>O<sub>6</sub>) C, H, N.

15

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Cyclooxygenase Inhibition Studies. The ability of the test compounds listed in Table 1 to inhibit ovine COX-1 and COX-2 (IC<sub>50</sub> value, μM) was determined using an enzyme immuno assay (EIA) kit (catalog no. 560101, Cayman Chemical, Ann Arbor, MI, USA) according to our previously reported method.<sup>38</sup>

25

Anti-inflammatory Assay. The test compounds 11-16 and the reference drugs (aspirin, ibuprofen and indomethacin) were evaluated using the in vivo rat carrageenan-induced foot paw edema model reported previously.<sup>39,40</sup>

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Nitric Oxide Release Assay: In vitro nitric oxide release, upon incubation with phosphate buffer, pig liver esterase, or guinea pig serum, was determined by quantification of nitrite produced by the reaction of nitric oxide with oxygen and water using the Griess reaction. Nitric oxide release data were acquired for test compounds (11-16), and the reference compounds O<sup>2</sup>-sodium 1-(pyrrolidin-1-yl)diazene-1,2-diolate, and O<sup>2</sup>-sodium 1-(N,N-dimethylamino)diazene-1,2-diolate (7) using the reported procedures.<sup>41</sup>

10

Acute Ulcerogenesis Assay: The ability to produce gastric damage was evaluated according to a reported procedure.<sup>42</sup> Ulcerogenic activity was evaluated after oral administration of aspirin (250 mg/kg), ibuprofen (250 mg/kg), indomethacin (30 mg/kg) or an equivalent amount of the correspondent test compound (11-16). All drugs were suspended and administered in 1.7 mL of a 1% methylcellulose solution. Control rats received oral administration of vehicle (1.7 mL of 1.0% methylcellulose solution). Food, but not water, was removed 24 h before administration of test compounds. Six hours after oral administration of the drug, rats were euthanized in a CO<sub>2</sub> chamber and their stomachs were removed, cut out along the greater curvature of the stomach, gently rinsed with water and placed on ice. The number and the length of ulcers were determined using a magnifier lens. The severity of the gastric lesion was measured along its greatest length (1 mm = rating of 1, 1-2 mm = rating of 2, >2 mm = rating according to their length in mm). The average overall length (in mm) of individual ulcers in each tissue was designated as the "ulcer index". Each experimental group consisted of four rats.

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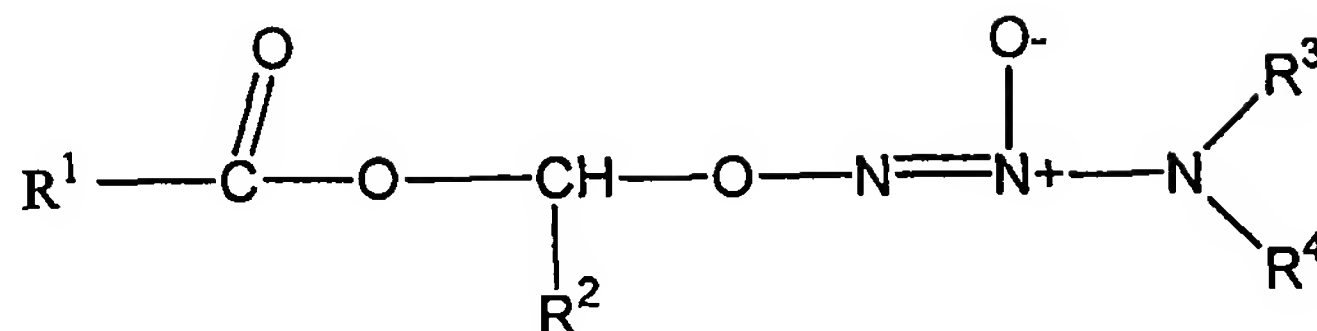


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What is claimed is:

1. A compound of the formula :



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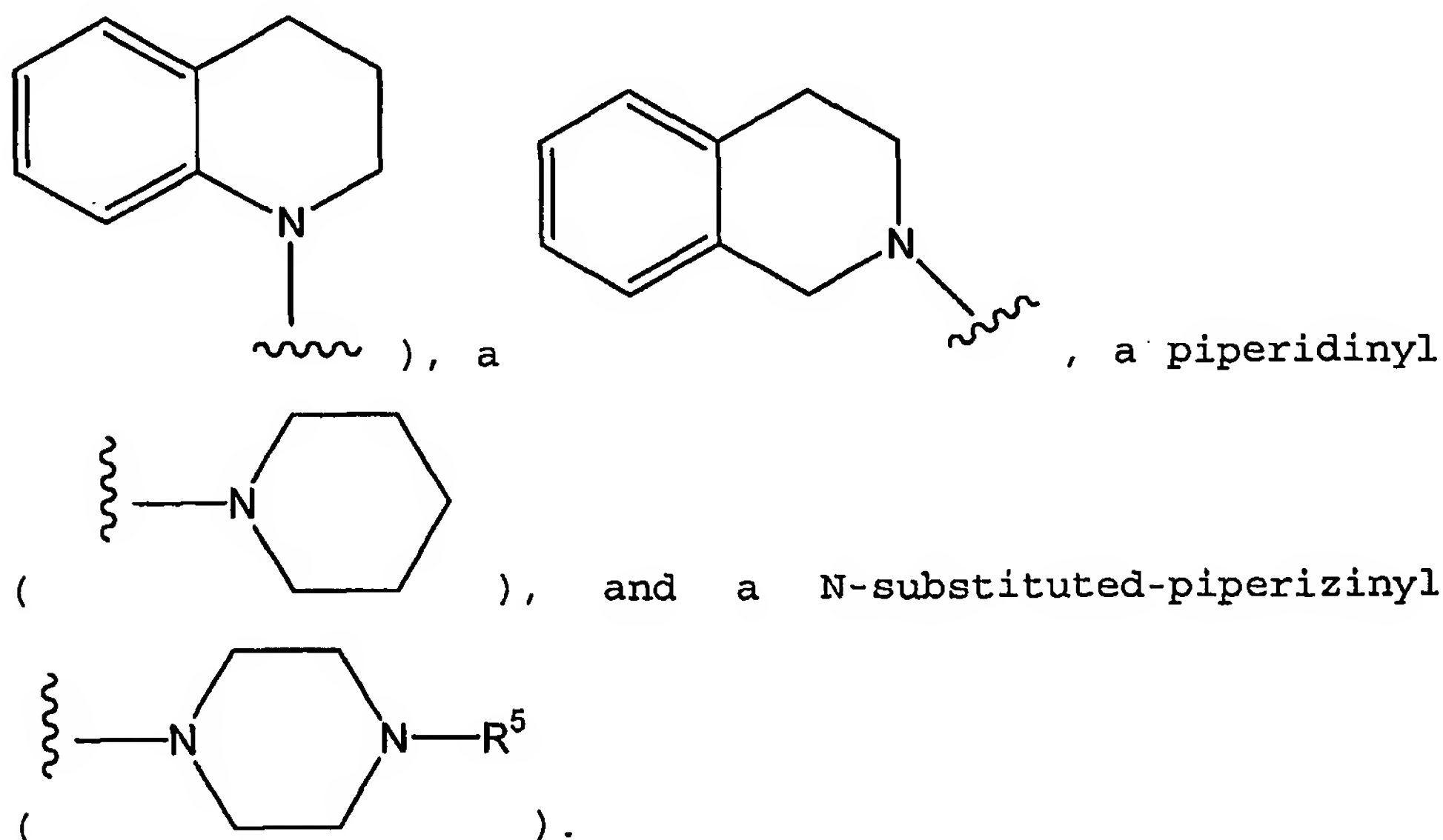
wherein R<sup>1</sup> is an uncarboxylated core of a non-steroidal anti-inflammatory drug,

10 R<sup>2</sup> is selected from the group consisting of a hydrogen, a C<sub>1-12</sub> straight chain alkyl, a C<sub>3-12</sub> branched chain alkyl, a C<sub>3-12</sub> straight chain alkenyl, a C<sub>3-12</sub> branched chain alkenyl, a C<sub>3-8</sub> cycloalkyl, an alkoxy, a nitrile, a halo, a morpholino, an amino, a benzyl, a phenyl, a C<sub>1-4</sub> aryl alkyl, a heteroaryl, an arylamino, a dialkylamino, a diarylamino, a carboxyalkylamino, a carboxydialkylamino, a tolyl, a xylyl, an anisyl, a mesityl, an acetoxyl, a carboxyl, a carboxyethyl, an alkylcarbonyl, a thiol, an alkylthiol, an alkyloxy, a  
15 carboxyamido, an alkylcarboxyamido, a dialkylcarboxyamido, a phenoxy, a benzyloxy, a phenylcarbonyl, a benzylcarbonyl, a nitrophenyl, a trialkylsilyl and a nitro;

20 R<sup>3</sup> and R<sup>4</sup> are selected from the group consisting of C<sub>1-12</sub> straight chain alkyl, a C<sub>3-12</sub> branched chain alkyl, a C<sub>3-12</sub> straight chain alkenyl, a C<sub>3-12</sub> branched chain alkenyl, a C<sub>3-8</sub> cycloalkyl, a morpholino, an amino, a benzyl, a C<sub>1-4</sub> aryl alkyl.

- 25  
30 2. The compound of claim 1, wherein R<sup>3</sup> and R<sup>4</sup> are same or different from each other.

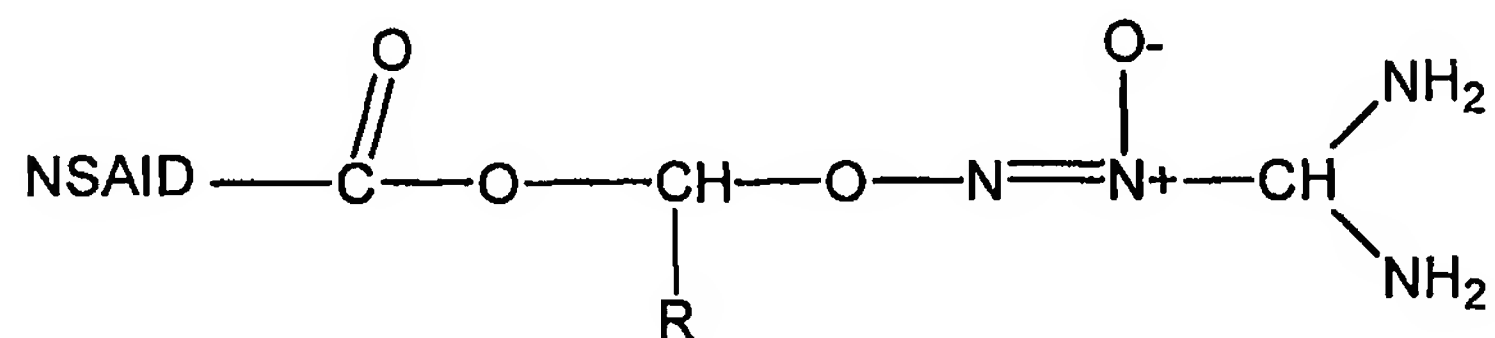
3. The compound of claim 1, wherein the  $R^2$  group is substituted or unsubstituted.
- 5 4. The compound of claim 1, wherein the  $-N(R^3, R^4)$  group is cyclized to form a structure selected from the group consisting of a 1,2,3,4-tetrahydroquinolyl (



5. The compound of claim 4, wherein  $R^5$  is selected from the group consisting of a  $C_{1-12}$  straight chain alkyl, a  $C_{3-12}$  branched chain alkyl, a  $C_{3-12}$  straight chain alkenyl, a  $C_{3-12}$  branched chain alkenyl, a  $C_{3-8}$  cycloalkyl, a benzyl, a phenyl, a  $C_{1-4}$  aryl alkyl, a heteroaryl, a tolyl, a xylyl, an anisyl, a mesityl, a carboxyethyl, an alkylcarbonyl, a phenylcarbonyl, a benzylcarbonyl, a nitrophenyl, and a trialkylsilyl.
- 20 6. The compound of claim 5, wherein the  $R^5$  group is substituted or unsubstituted.

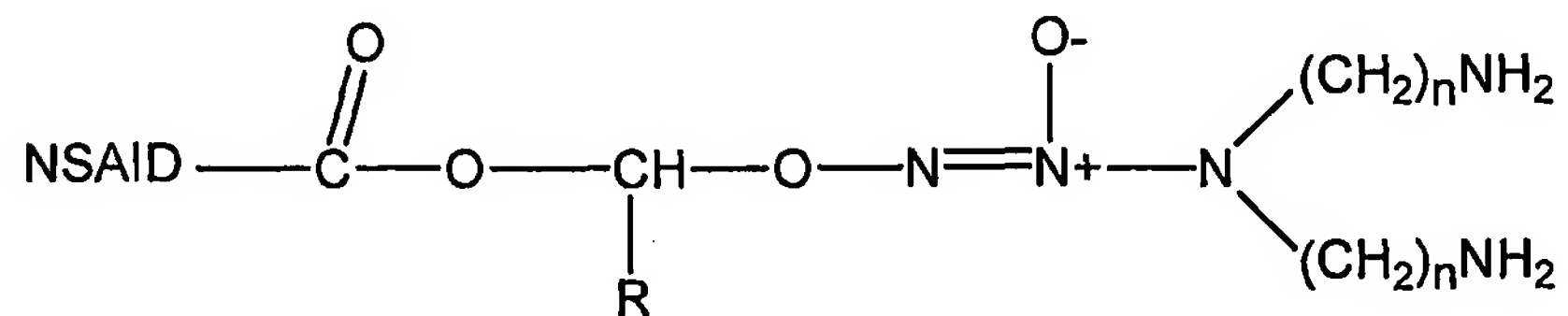
7. The compound of claim 1, wherein the non-steroidal anti-inflammatory drug is selected from the group consisting of acetylsalicylic acid, ibuprofen, naproxen, indomethacin, salicylic acid, diflunisal, salsalate, olsalazine, sulfasalazine, sulindac, etodolac, mefenamic acid, meclofenamic acid, tolmetin, ketorolac, diclofenac, fenoprofen, ketoprofen, oxaprozin, carprofen, flurbiprofen, nabumetone, other related carboxylic acids with anti-inflammatory activity, and their pharmaceutically suitable salts.

8. A compound of the formula :



- wherein R is same as the R<sub>2</sub> in the compound of Claim 1, and the compound includes pharmaceutically suitable alkali metal salts or hydrochloride salts thereof.

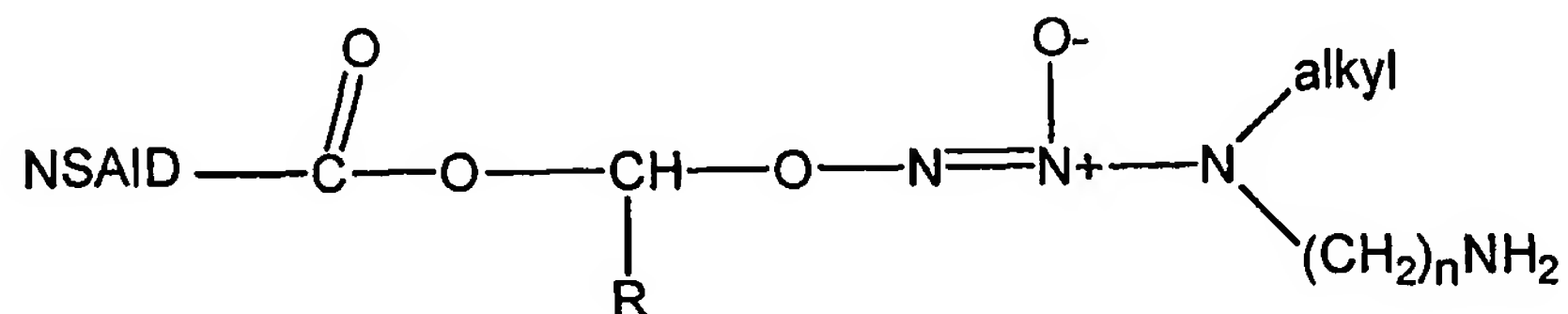
9. A compound of the formula :



wherein R is same as the R<sub>2</sub> in the compound of Claim 1, n=1-8, and the compound includes pharmaceutically suitable alkali metal salts or hydrochloride salts thereof.

5

10. A compound of the formula:

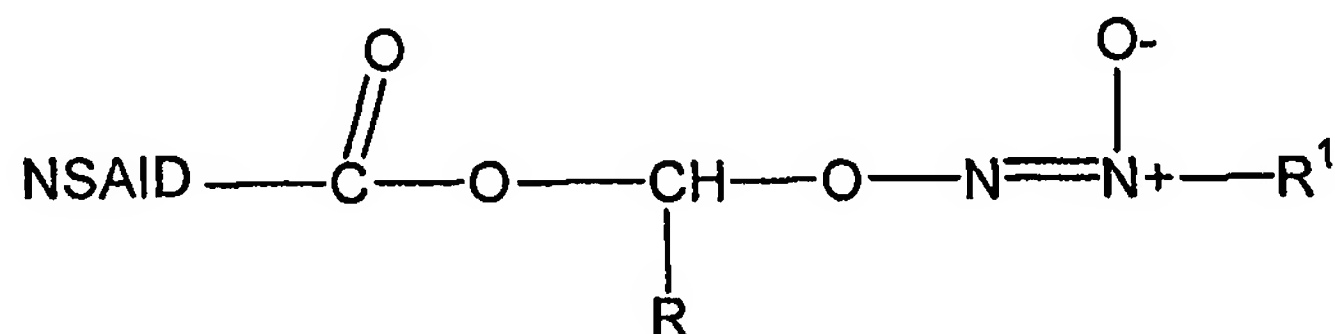


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wherein R is same as the R<sub>2</sub> in the compound of Claim 1, n=1-8, and the compound includes pharmaceutically suitable alkali metal salts or hydrochloride salts thereof.

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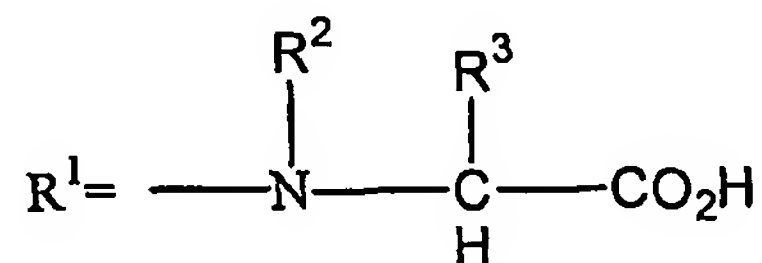
11. A compound of the formula:



20

wherein R is same as the R<sub>2</sub> in the compound of Claim 1, and R<sub>1</sub> is a N-substituted amino acid moiety.

12. The compound of claim 11, wherein the N-substituted amino acid moiety is:



25

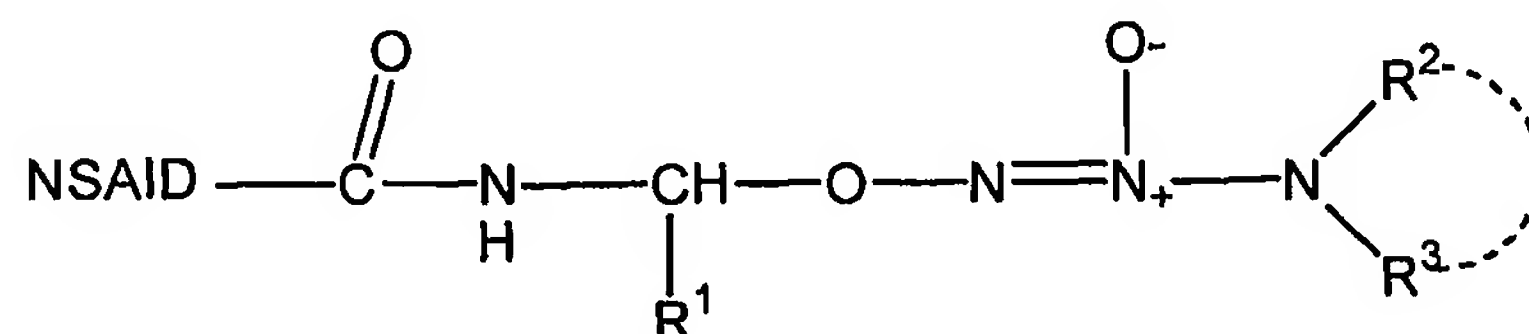
wherein R<sub>2</sub> is selected from the group consisting of a hydrogen, a C1-12 straight chain alkyl, a C3-12

branched chain alkyl, a C3-12 straight chain alkenyl, a  
 C3-12 branched chain alkenyl, a C3-8 cycloalkyl, a  
 benzyl, a phenyl, a C1-4 aryl alkyl, a heteroaryl, a  
 tolyl, a xylyl, an anisyl, a mesityl, a carboxyethyl,  
 5 and R3 is selected from the group consisting of a  
 hydrogen, a C1-12 straight chain alkyl, a C3-12  
 branched chain alkyl, a C3-12 straight chain alkenyl, a  
 C3-12 branched chain alkenyl, a C3-8 cycloalkyl, an  
 alkoxy, a nitrile, a halo, a morpholino, an amino, a  
 10 benzyl, a phenyl, a C1-4 aryl alkyl, a heteroaryl, an  
 arylamino, a dialkylamino, a diarylamino, a  
 carboxyalkylamino, a carboxydialkylamino, a tolyl, a  
 xylyl, an anisyl, a mesityl, an acetoxy, a carboxy, a  
 carboxyethyl, an alkylcarbonyl, a thiol, an alkylthiol,  
 15 an alkyloxy, a carboxyamido, an alkylcarboxyamido, a  
 dialkylcarboxyamido, a phenoxy, a benzyloxy, a  
 phenylcarbonyl, a benzylcarbonyl, a nitrophenyl, a  
 trialkylsilyl, and a nitro.

20 13. The compound of claim 12, wherein the N-substituted amino  
 acid moiety is selected from the group consisting of N-  
 methylglycine, N-methylalanine, N-methylphenylalanine, N-  
 methylserine, and any other N-alkyl amino acid.

25 14. The compound of claim 12, wherein the R2 and R3 are  
 substituted or unsubstituted.

15. An amide bioisostere ester compound of the formula:

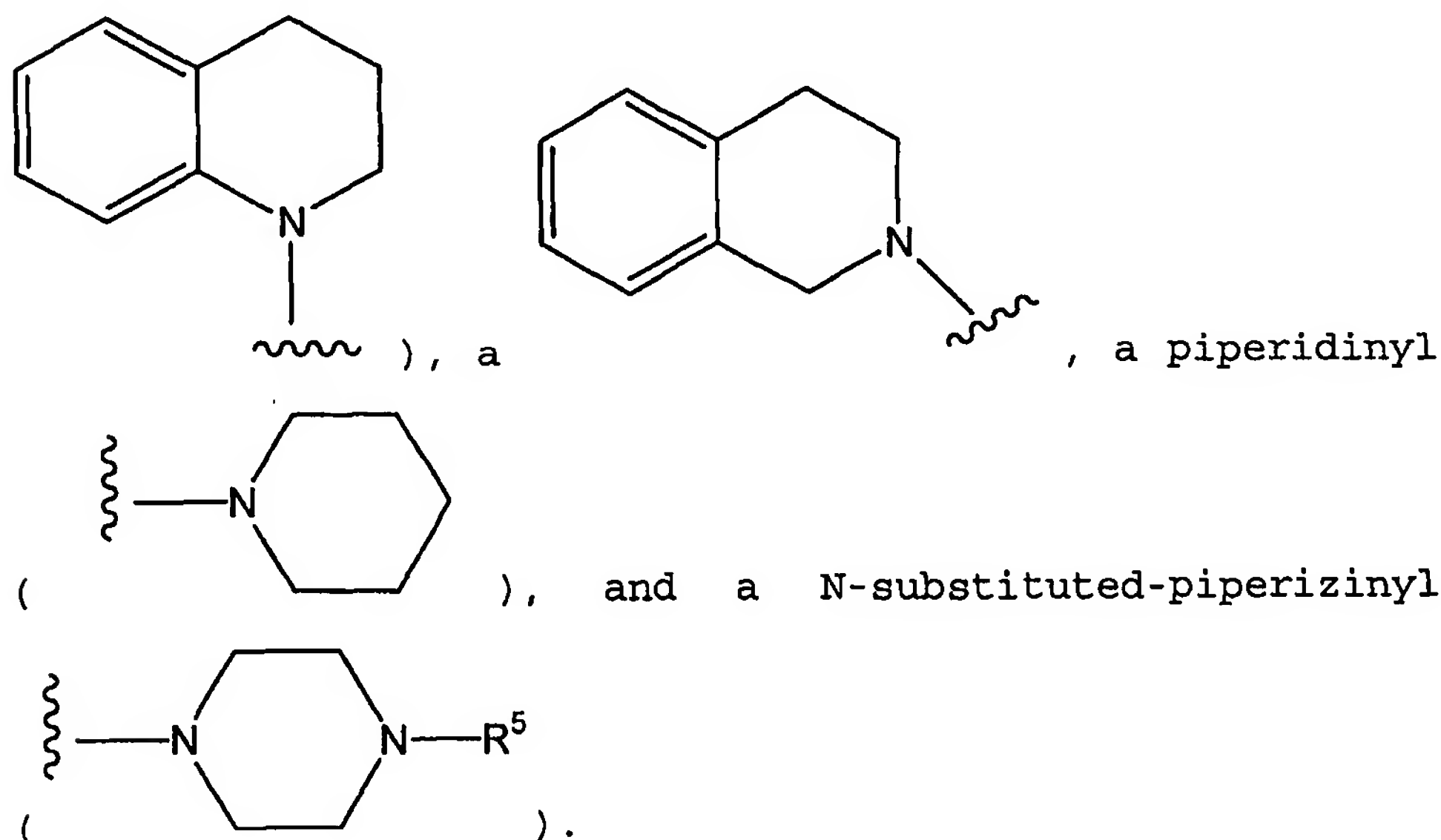


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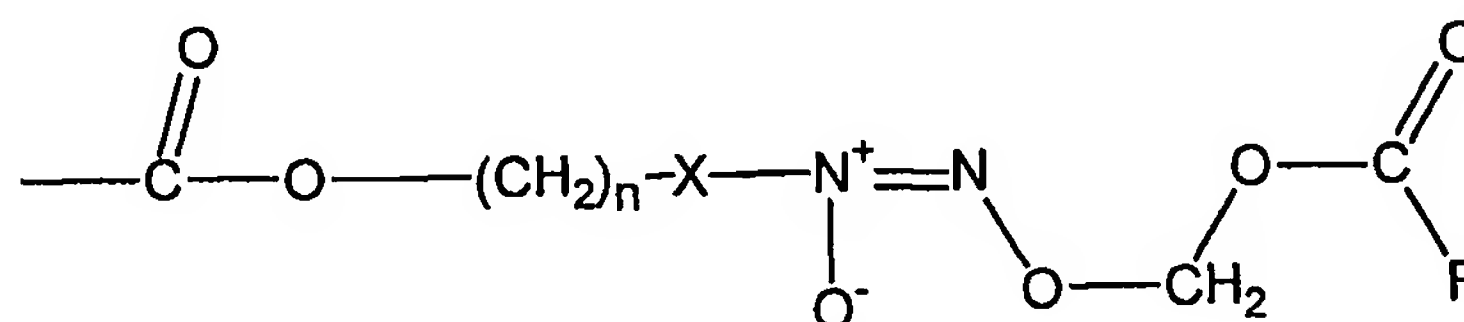


## Structure X

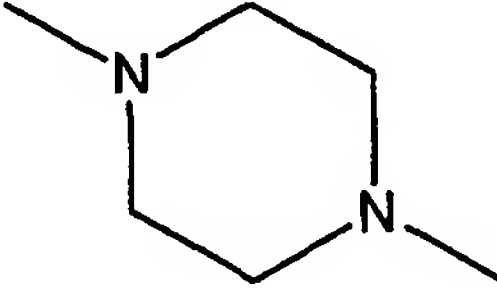
wherein R1 is selected from the group consisting of a  
hydrogen, a C1-12 straight chain alkyl, a C3-12  
5 branched chain alkyl, a C3-12 straight chain alkenyl, a  
C3-12 branched chain alkenyl, a C3-8 cycloalkyl, an  
alkoxy, a nitrile, a halo, a morpholino, an amino, a  
benzyl, a phenyl, a C1-4 aryl alkyl, a heteroaryl, an  
arylamino, a dialkylamino, a diarylamino, a  
10 carboxyalkylamino, a carboxydialkylamino, a tolyl, a  
xylyl, an anisyl, a mesityl, an acetoxy, a carboxy, a  
carboxyethyl, an alkylcarbonyl, a thiol, an alkylthiol,  
an alkyloxy, a carboxyamido, an alkylcarboxyamido, a  
dialkylcarboxyamido, a phenoxy, a benzyloxy, a  
15 phenylcarbonyl, a benzylcarbonyl, a nitrophenyl, a  
trialkylsilyl, and a nitro, and  
the -N(R2, R3) group is cyclized to form a structure  
selected from the group consisting of a 1,2,3,4-  
tetrahydroquinolyl (

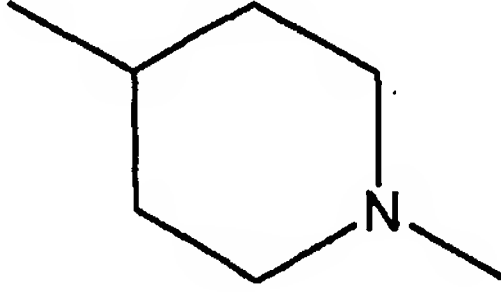


16. The compound of claim 15, wherein R<sup>5</sup> is selected from the group consisting of a C<sub>1-12</sub> straight chain alkyl, a C<sub>3-12</sub> branched chain alkyl, a C<sub>3-12</sub> straight chain alkenyl, a C<sub>3-12</sub> branched chain alkenyl, a C<sub>3-8</sub> cycloalkyl, a benzyl, a phenyl, a C<sub>1-4</sub> aryl alkyl, a heteroaryl, a tolyl, a xylyl, an anisyl, a mesityl, a carboxyethyl, an alkylcarbonyl, a phenylcarbonyl, a benzylcarbonyl, a nitrophenyl, and a trialkylsilyl.
17. The compound of claim 16, wherein the R<sup>5</sup> group is substituted or unsubstituted.
18. The compound of claim 15, wherein the R<sup>1</sup> group is substituted or unsubstituted.
19. A compound of the formula:



wherein X is selected from the group consisting of N-

substituted piperiziny (  ), N-

and 4-substituted piperidinyl (  ), and a N-methyl moiety, and

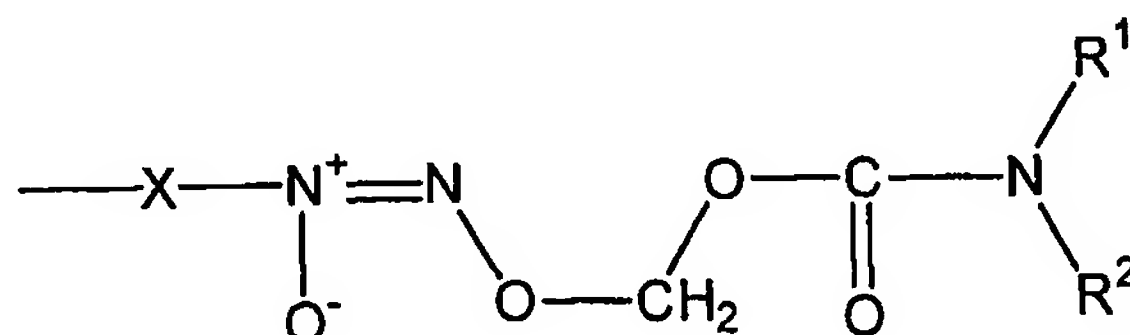
R is selected from the group consisting of a C<sub>1-12</sub> straight chain alkyl, a C<sub>3-12</sub> branched chain alkyl, a C<sub>3-12</sub> straight chain alkenyl, a C<sub>3-12</sub> branched chain alkenyl, a C<sub>3-8</sub> cycloalkyl, an alkoxy, a nitrile, a halo, a morpholino, an amino, a benzyl, a phenyl, a C<sub>1-4</sub> aryl alkyl, a heteroaryl, an arylamino, a dialkylamino, a diarylamino, a carboxyalkylamino, a carboxydialkylamino, a tolyl, a xylyl, an anisyl, a mesityl, an acetoxyl, a carboxyl, a carboxyethyl, an alkylcarbonyl, an alkylthiol, an alkyloxy, a carboxyamido, an alkylcarboxyamido, a dialkylcarboxyamido, a phenoxy, a benzyloxy, a phenylcarbonyl, a benzylcarbonyl, a nitrophenyl, a trialkylsilyl, and a nitro.

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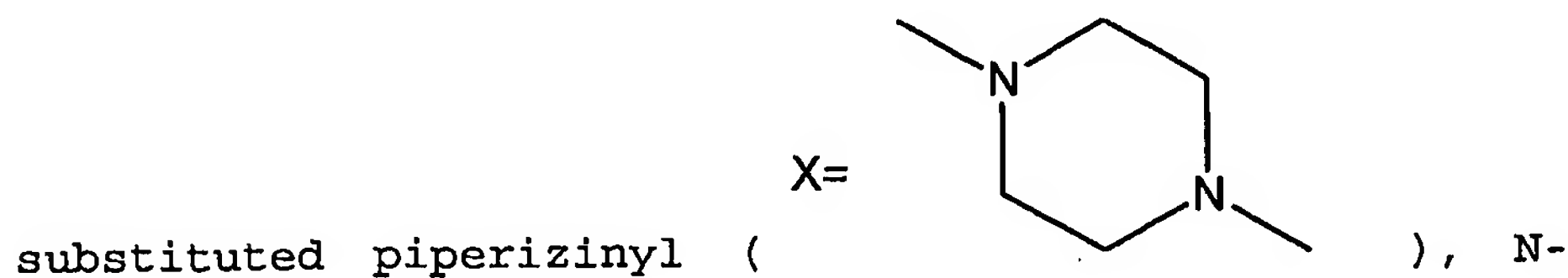
20. The compound of claim 19, wherein the R group is substituted or unsubstituted.

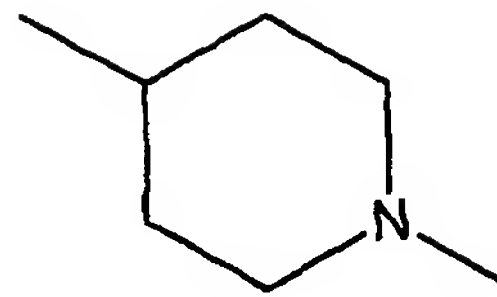
21. A carbamate compound of the formula:

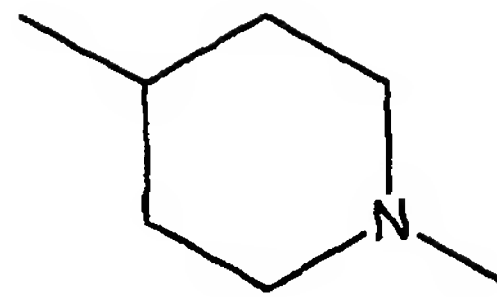
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wherein X is selected from the group consisting of N-





and 4-substituted piperidinyl (  ), and a N-methyl moiety, and

$R^1$  and  $R^2$  are selected from the group consisting of a hydrogen, a  $C_{1-12}$  straight chain alkyl, a  $C_{3-12}$  branched chain alkyl, a  $C_{3-12}$  straight chain alkenyl, a  $C_{3-12}$  branched chain alkenyl, a  $C_{3-8}$  cycloalkyl, a benzyl, a phenyl, a  $C_{1-4}$  aryl alkyl, a heteroaryl, a tolyl, a xylyl, an anisyl, a mesityl, a carboxyethyl, an alkylcarbonyl, a phenylcarbonyl, a benzylcarbonyl, a nitrophenyl, a trialkylsilyl, and a nitro.

22. The compound of claim 21, wherein  $R_1$  and  $R_2$  are substituted or unsubstituted.

23. A compound  $O^2$ -(Acetylsalicyloyloxymethyl)-1-(pyrrolidin-1-yl)diazen-1-ium-1,2-diolate as shown in Figure 6.

24. A compound  $O^2$ -(Acetylsalicyloyloxymethyl)-1-(N,N-dimethylamino)diazen-1-ium-1,2-diolate as shown in Figure 6.

25. A compound  $O^2$ -[2-(4-(Isobutyl)phenyl)propanoyloxymethyl]-1-(pyrrolidin-1-yl)diazen-1-ium-1,2-diolate as shown in Figure 6.

26. A compound  $O^2$ -[2-(4-(Isobutyl)phenyl)propanoyloxymethyl]-1-(N,N-dimethylamino)diazen-1-ium-1,2-diolate as shown in Figure 6.

27. A compound  $O^2$ -[2-(1-(4-Chlorobenzoyl)-5-methoxy-2-methyl-1H-indol-3-yl)acetoxymethyl]-1-(pyrrolidin-1-yl)diazen-1-ium-1,2-diolate as shown in Figure 6.
- 5 28. A compound  $O^2$ -[2-(1-(4-Chlorobenzoyl)-5-methoxy-2-methyl-1H-indol-3-yl)acetoxymethyl]-1-(dimethyl amino)diazen-1-ium-1,2-diolate as shown in Figure 6.
- 10 29. A composition comprising an effective amount of one of the compounds of claims 1-22 in the same molar dose range as recommended for the NSAID from which it was derived.
- 15 30. A composition comprising an effective amount of one of the compounds of claims 1-22 in various dose ranges capable of enhancing therapeutic outcome as recommended for the NSAID from which it was derived.
- 20 31. Use of one of the compounds of claims 1-22 for reducing gastrointestinal side effects of a parent non-steroidal anti-inflammatory drugs in a subject.
- 25 32. Use of one of the compounds of claims 1-22 in the manufacture of medicament for reducing gastrointestinal side effects of a parent non-steroidal anti-inflammatory drugs in a subject.
- 30 33. The use of claim 31 or 32, wherein the side effects are selected from the group consisting of dyspepsia, nausea, vomiting, abdominal pain, diarrhea, gastric bleeding, intestinal bleeding, gastric ulceration, and intestinal ulceration.

34. Use of one of the compounds of Claims 1-22 for the indications recommended for the unsubstituted NSAID from which it is derived.
- 5 35. Use of one of the compounds of Claims 1-22 in the manufacture of medicament for the indications recommended for the unsubstituted NSAID from which it is derived.
- 10 36. The use of Claim 34 or 35, wherein the indication is selected from the group consisting of pain, inflammation, and headache.
37. The use of Claim 36, wherein the unsubstituted NSAID is ibuprofen.
- 15 38. The use of Claim 34 or 35, wherein the indication is cardiovascular protection.
- 20 39. The use of Claim 38, wherein the unsubstituted NSAID is acetylsalicylic acid.
40. The use of Claim 34 or 35, wherein the indication is rheumatoid or osteoarthritis symptoms.
- 25 41. The use of Claim 40, wherein the unsubstituted NSAID is naproxen or indomethacin

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Figure 1-1

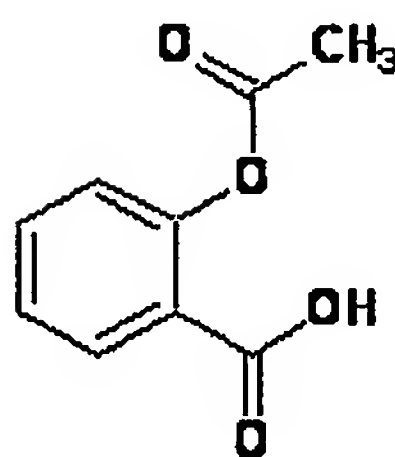


Figure 1-2

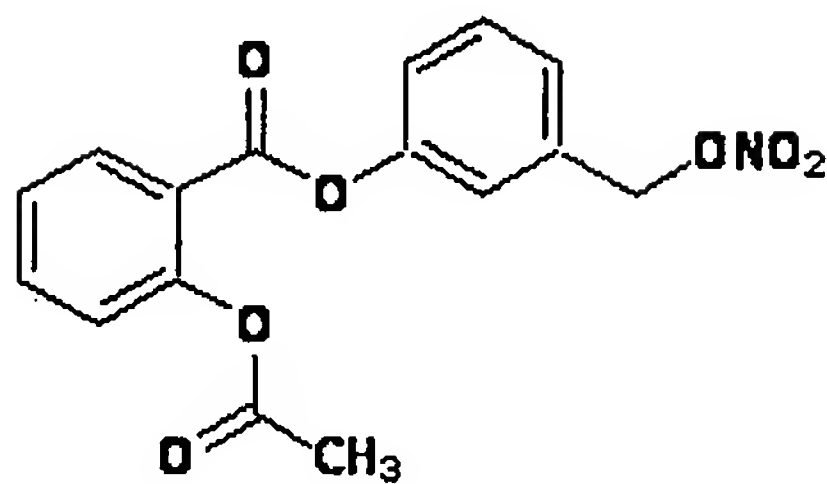
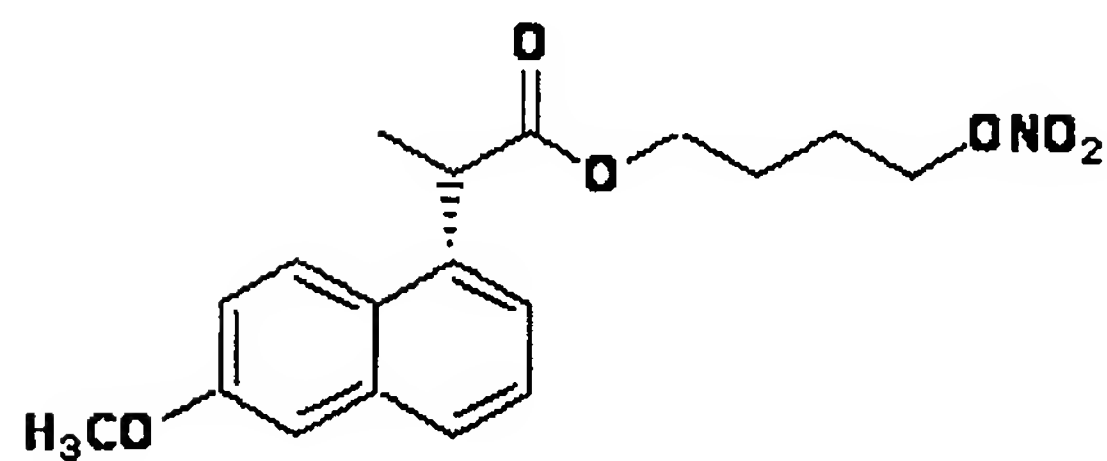


Figure 1-3





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Figure 1-4

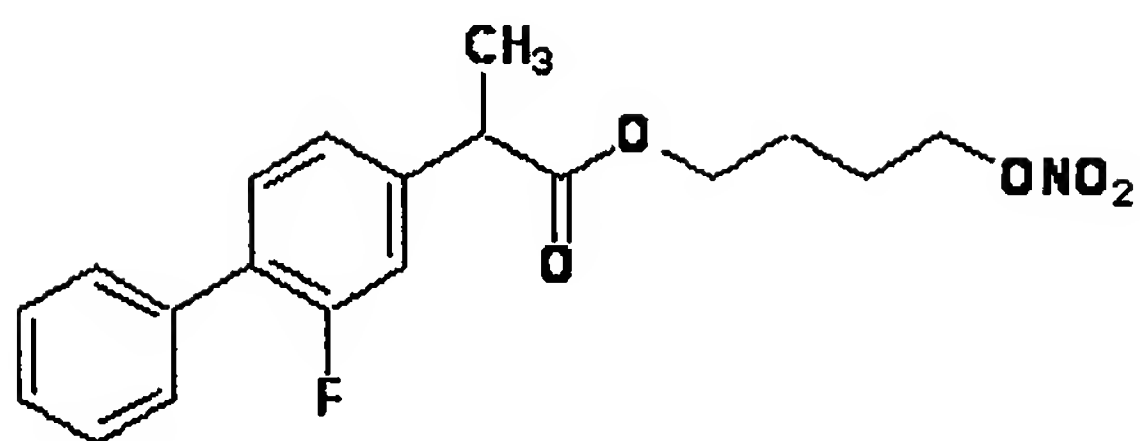
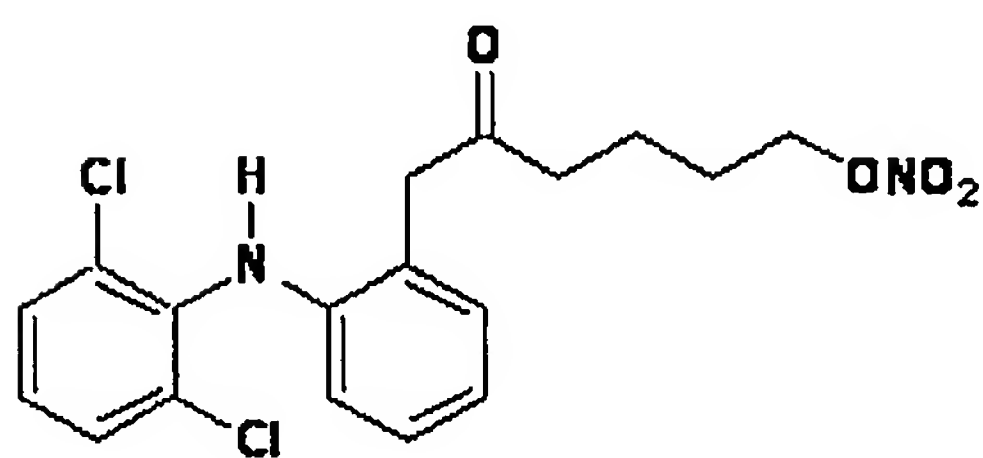
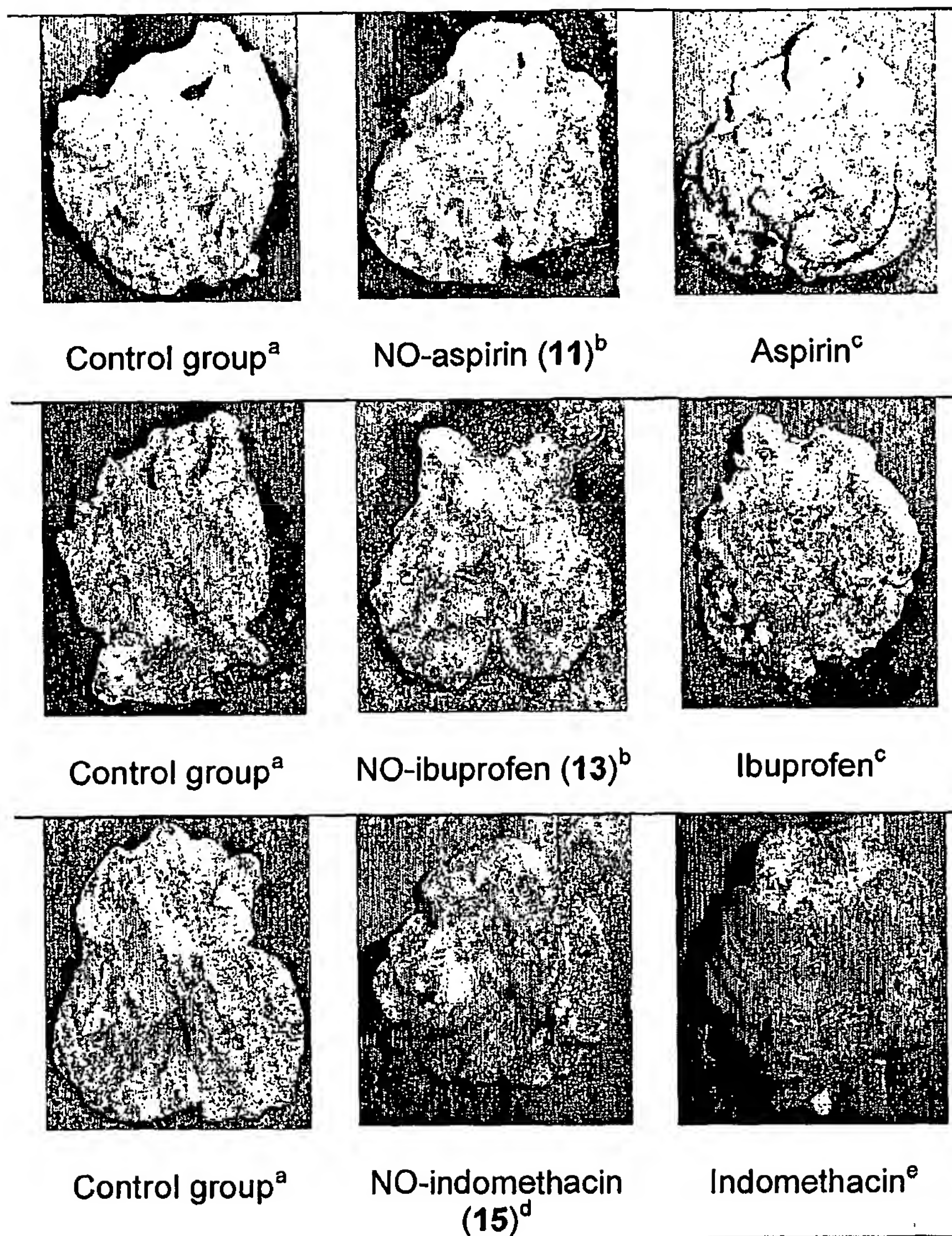


Figure 1-5



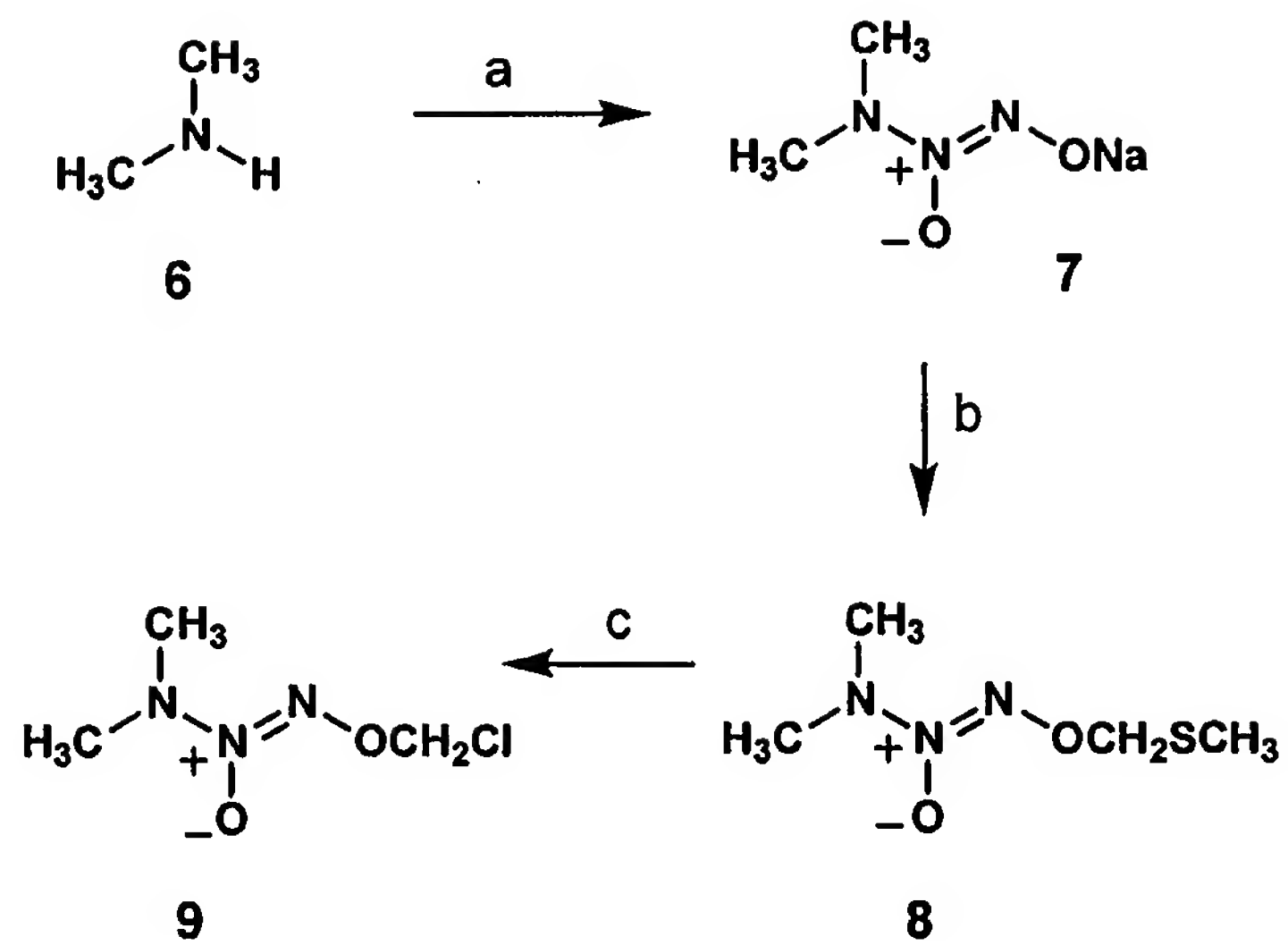
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Figure 2



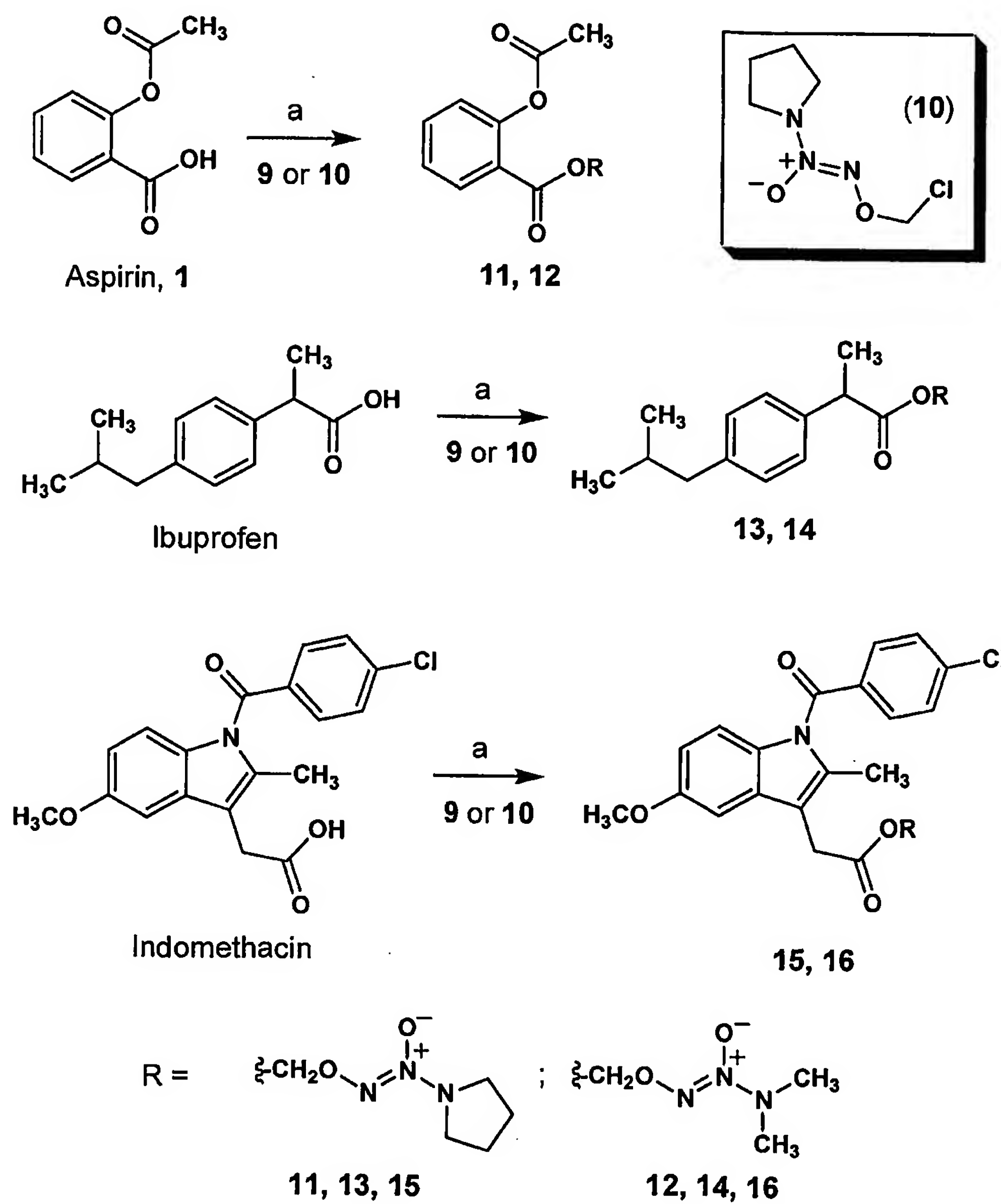
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Figure 3



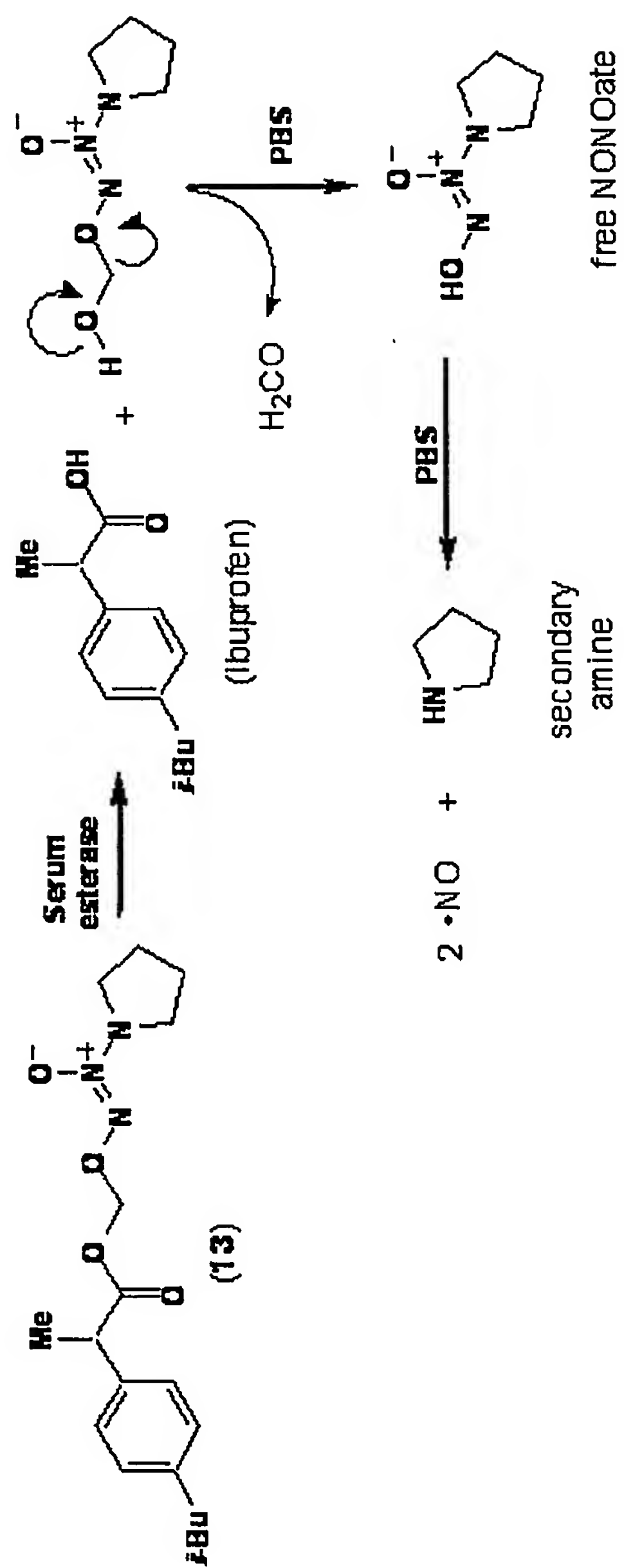
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Figure 4.



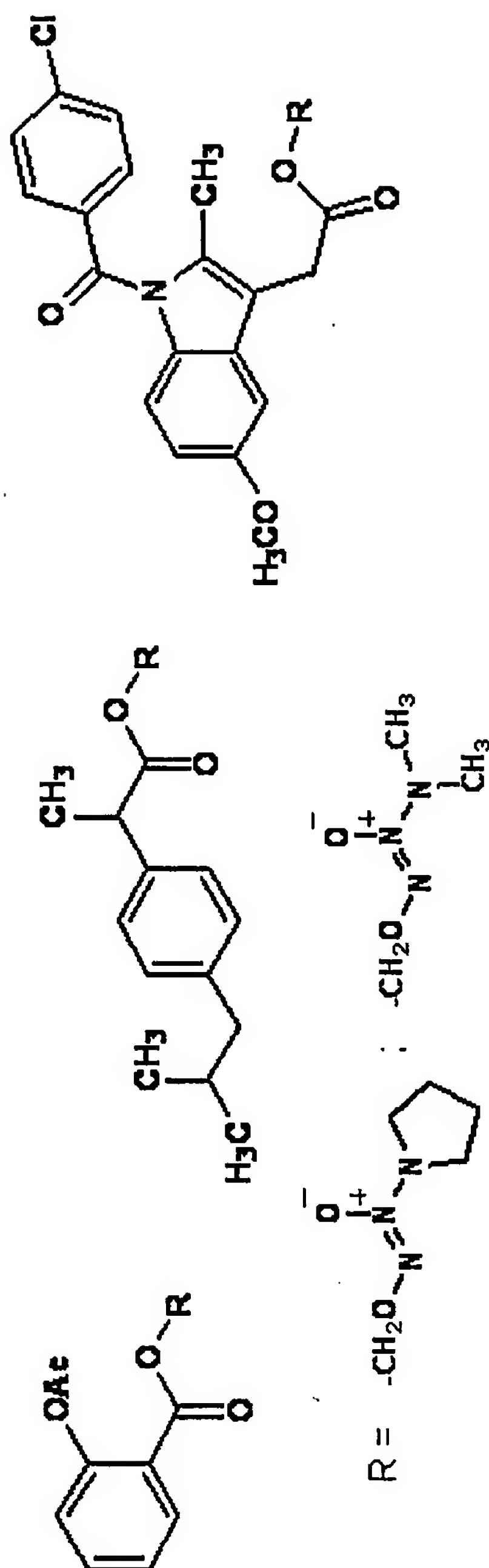
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Figure 5



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Figure 6



# INTERNATIONAL SEARCH REPORT

International application No.

PCT/US06/19115

## A. CLASSIFICATION OF SUBJECT MATTER

IPC(8): A61K 31/655( 2006.01);C07C 245/24( 2006.01)

USPC: 514/149;534/550

According to International Patent Classification (IPC) or to both national classification and IPC

## B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 514/149; 534/550

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)  
STN CAS ON LINE

## C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US 6,949,530 B2 (HARBIE et al) 27 September 2005 (27.09.2005) entire document	1-30

☐

Further documents are listed in the continuation of Box C.

☐

See patent family annex.

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Date of the actual completion of the international search

16 August 2006 (16.08.2006)

Date of mailing of the international search report

15 SEP 2006

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Mail Stop PCT, Attn: ISA/US  
Commissioner for Patents  
P.O. Box 1450  
Alexandria, Virginia 22313-1450

Facsimile No. (571) 273-3201

Authorized officer

Kamal A. Saeed

Telephone No. (571) 272 1600